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DEVELOPMENT OF A BIPOLAR LEAD/ACID
BATTERY FOR THE MORE ELECTRIC AIRCRAFT



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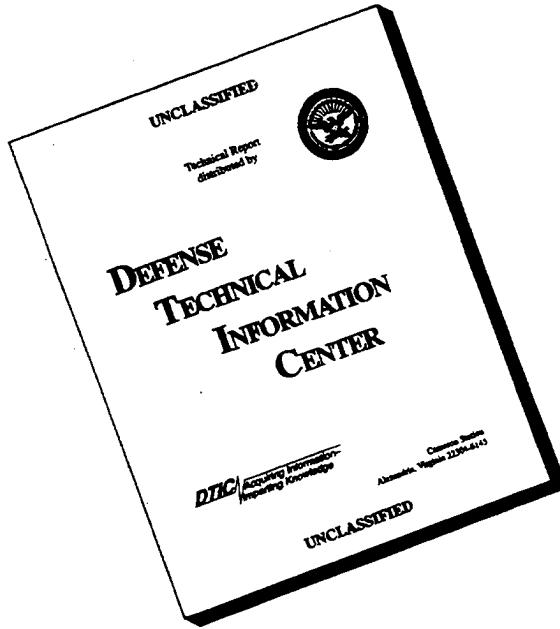
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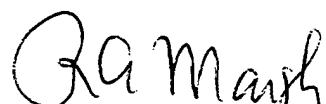
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This report summarizes the development work completed under contract F33615-91-C-2142 for the time period of September 1991 to September 1995. Initial work targeted the development of a filled polymeric composite substrate for use in a true bipolar lead acid battery. Efforts were refocused on metallic substrate technology in Month 33, and concluded with the delivery of battery systems to Wright Laboratory.				
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1.0 SUMMARY

A 36-month contract was undertaken by Johnson Controls Battery Group, Inc. to develop a highly conductive, non-porous, and lightweight bipolar substrate and deliver a 56-volt prototype module for evaluation for the More Electric Aircraft. Eighteen months into Contract #F33615-91-C-2142, significant accomplishments were reported in the identification of suitable composite materials and in optimizing the compounding parameters of same. Laminated, 8 cm (L) x 8 cm (H) x 0.102 cm (TH) substrates with an overall resistivity of 4-6 Ω -cm were routinely manufactured in-house and used in battery builds. Over 150 cycles were demonstrated to 100% DOD at 0.16 A/cm² in a 4-volt battery configuration. Mass production oriented container molding was also demonstrated, however, process reliability was a major concern. Critical evaluation of the project in Month 33 recognized the difficulties in addressing recurrent substrate and paste adhesion delamination, as well as those to be solved in achieving high power (0.48+ A/cm²) capability from a 400+ cm² electrode. High power capability from a composite substrate was not deemed likely in the remaining contract period. Therefore, given its success in a parallel internally funded bipolar program, JCBGI requested a no-cost time extension to evaluate a new approach in metallic bipolar substrate technology. Five attempts were made at cladding strips of various corrosion resistant alloys, however, resultant materials were never suited to pasting or battery builds. Concurrent efforts to redesign the injection molded container succeeded in eliminating internal distortion of the metallic electrodes, but failed to resolve cell-to-cell leakage around the fill ports. At contract's end, deliverables utilizing a binary lead alloy and an alternative containment design were assembled, formed and delivered to WPAFB for test and evaluation.

Future composite bipolar substrate investigations based upon this body of work should focus on fostering positive paste adhesion. Continued metallic substrate work would benefit most from efforts to increase the substrate strength and corrosion resistance. Both designs require additional development of the injection molded containment concept to eliminate the catastrophic cell-to-cell leakage exhibited at the close of this contract.

2.0. WORK BREAKDOWN SCHEDULE

As with other contract work performed at JCBGI, a Work Breakdown Schedule (WBS) was implemented to plan, execute, and monitor technical progress, costs, and scheduling. Tasks were identified as unitary efforts necessary to complete individual aspects of battery development, and subtasks further delineated each task. Composite plans, shown in Figure 1, were easily translated in August 1994 to more closely describe the efforts necessary to assemble a 24-volt bipolar battery utilizing metallic based substrates. These interpretations are shown in parentheses next to the composite substrate counterparts within Figure 1.

FIGURE 1: BMET WORK BREAKDOWN SCHEDULE

WBS 1.0 PROGRAM MANAGEMENT

- Subtask 1.1 Managing Strategy
- Subtask 1.2 Liaison/Meetings
- Subtask 1.3 Documentation
- Subtask 1.4 Contract Administration
- Subtask 1.5 Operating Supplies

WBS 2.0 BATTERY DESIGN

- Subtask 2.1 Battery System Design Analysis
- Subtask 2.2 Performance Modeling

WBS 3.0 BIPOLAR PLATE

- Subtask 3.1 Conductive Fillers (Multi-Alloy Substrate Development)
- Subtask 3.2 Substrate Fabrication Processes (Rolling/Embossing Work)
- Subtask 3.3 Stability Testing (Corrosion Testing)
- Subtask 3.4 Proof of Concept Testing (Small Scale Characterization)

WBS 4.0 BATTERY COMPONENTS

- Subtask 4.1 Separator Material
- Subtask 4.2 Active Material Development (Freeze/Thaw Work)

WBS 5.0 BATTERY FABRICATION

- Subtask 5.1 Sealing Methods (Lead to Plastic Interface Seal)
- Subtask 5.2 Formation

WBS 6.0 BMET DEMONSTRATION

- Subtask 6.1 Battery Fabrication (Deliverables)
- Subtask 6.2 Testing (Group 34 Cycling)

3.0 COMPOSITE SUBSTRATE DEVELOPMENT

3.1 WBS 1.0 PROGRAM MANAGEMENT

3.1.1 Subtask 1.1 Managing Strategy

Five review meetings were scheduled and attended by WPAFB and JCBGI personnel. These dates, as well as milestones achieved during the composite development phase of the contract, are shown in Gantt chart form in Figure 2.

3.2 WBS 2.0 BATTERY DESIGN

3.2.1 Subtask 2.1 Battery System Design Analysis

Preliminary performance requirements for the More Electric Aircraft (MEA) energy source were given to JCBGI by Richard Flake of WPAFB during the program kickoff meeting on December 12, 1991. The following energy sources were required:

Main Engine Starting:	150 kW, 30 sec
Ground Power:	25-75 kW, 30-45 min
Emergency Power:	75 kW, 10 min
APU Starting:	5-10 kW, 15 sec
Hybrid Emergency:	50-75 kW, 60 sec
Temperature Range:	-65°F to 120°F
Voltage Window:	270 volts (min), 330 volts (max)

Given this, JCBGI proceeded to use its proprietary lead/acid battery mathematical model to design near- and far-term bipolar systems having 5- and 10- year development time frames. Near-term modeling assumed that substrate program goals were reached and conventional active materials were used. The 10-year battery systems were projected assuming a thinner, more conductive substrate and improved active materials. The results, shown in Figures 3 through 14, dramatically illustrate the system configuration's dependence on application. Designs required as little as 0.18 ft³ with a system mass of 33 pounds to as much as 8.13 ft³ and 1349 pounds.

3.3 WBS 3.0 BIPOLAR PLATE

3.3.1 Subtask 3.1 Conductive Fillers

Initial work was focused on identifying an electronically conductive, filled polymeric composite having negligible ionic conduction which could short adjacent cells. The substrate was likewise required to be chemically inert to the electrode reactions, to have high oxygen and hydrogen overpotentials in H₂SO₄, and to be readily manufactured.

FIGURE 2: Composite Development Gantt Chart with Milestones

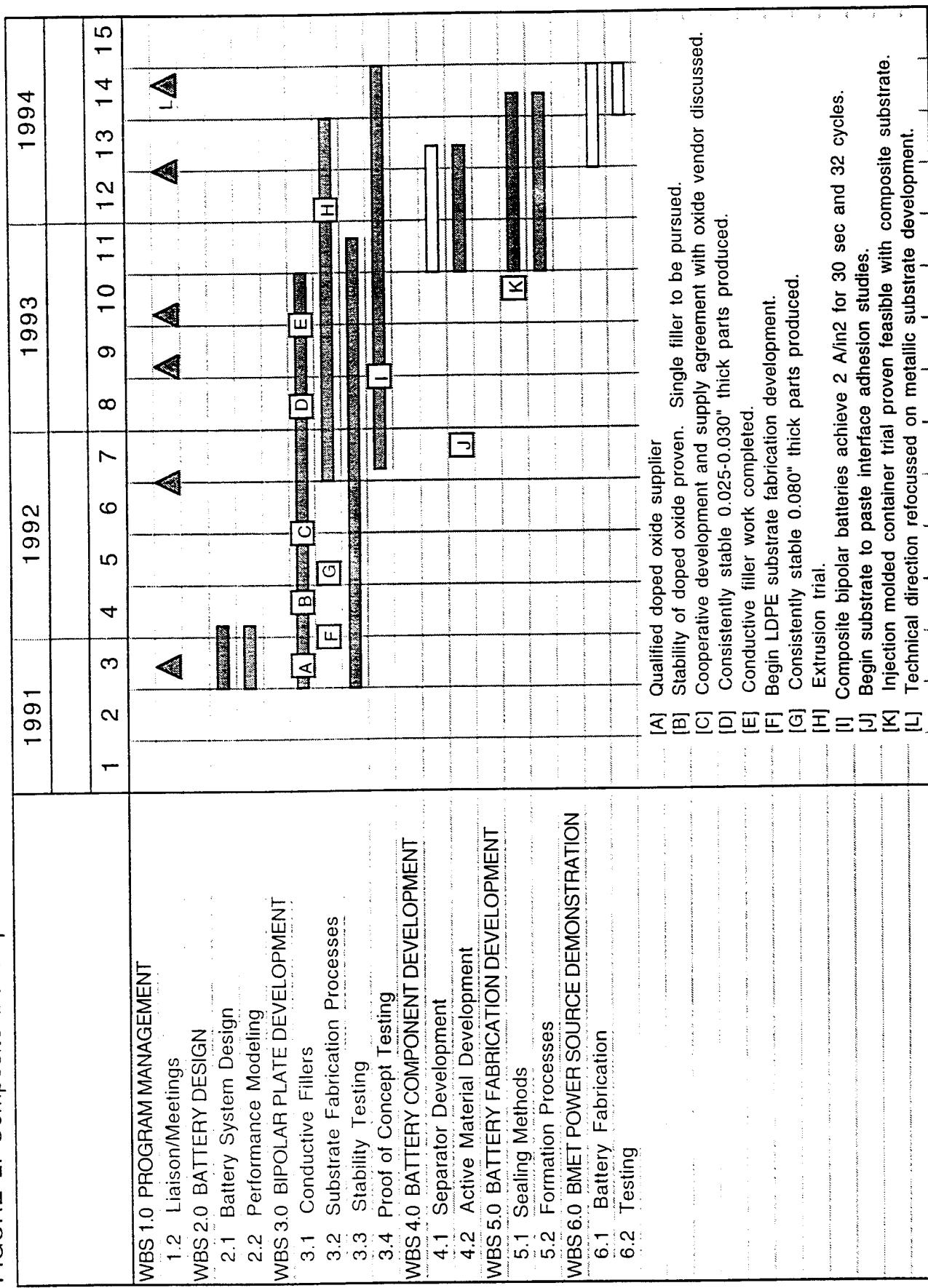


FIGURE 3

NEAR AND FAR TERM BMET BIPOLAR BATTERY SPECIFICATIONS

<u>BATTERY TYPE</u>	<u>NEAR TERM</u>	<u>FAR TERM</u>
---------------------	------------------	-----------------

Main Engine Starting

Mass	450 lbs.	389 lbs.
Volume	2.45 ft³	2.00 ft³

Ground Power

Lower Capacity Unit		
Mass	1000 lbs.	865 lbs.
Volume	6.15 ft³	4.85 ft³

Higher Capacity Unit		
Mass	1349 lbs.	1235 lbs
Volume	8.13 ft³	6.72 ft³

APU Starting

Mass Volume	33.4 lbs.	30.6 lbs
Volume	0.18 ft³	0.16 ft³

Assumptions:

Substrate Thickness	0.025"	0.010"
Substrate Weight	150 mg/cm²	80 mg/cm²
Substrate Resistivity	2.0 Ω-cm	~0 Ω-cm

FIGURE 4

BMET PERFORMANCE REQUIREMENTS
BIPOLAR BATTERY SPECIFICATIONS
 Near Term Projections (within 5 years)
 330 Volt Battery Systems

REQUIREMENTS MET	BATTERY DIMENSIONS	BATTERY VOLUME	BATTERY WEIGHT	W/kg	W/cm ³	W-hr/kg	W- hr/cm ³
Main Engine Starting APV Starting Hybrid Emergency	17.6"x15.5"x15.5"	2.45 ft ³	450 lbs	747.9	2.2	12.25	0.036
Main Engine Starting Ground Power Emergency Power APU Starting Hybrid Emergency							
Scenario 1 30 minute ground power capacity	27.4"x19.7"x19.7"	6.15 ft ³	1000 lbs	62.2	0.16	31.08	0.081
Scenario 2 45 minute ground power capacity	36.2"x19.7"x19.7"	8.13 ft ³	1349 lbs	46.1	0.12	34.56	0.092
APU Starting	16.5"x4.33"x4.33"	0.18 ft ³	33 lbs	705.0	2.1	11.75	0.036

FIGURE 5

BMET PERFORMANCE REQUIREMENTS
BIPOLAR BATTERY SPECIFICATIONS
Far Term Projections (10 years)
330 Volt Battery Systems

REQUIREMENTS MET	BATTERY DIMENSIONS	BATTERY VOLUME	BATTERY WEIGHT	W/kg	W/cm ³	W-hr/kg	W-hr/cm ³
Main Engine Starting APV Starting Hybrid Emergency	14.4"x15.5"x15.5"	2.00 ft ³	389 lbs	895.3	2.8	14.17	0.044
Main Engine Starting Ground Power Emergency Power APU Starting Hybrid Emergency							
Scenario 1 30 minute ground power capacity	21.6"x19.7"x19.7"	4.85 ft ³	864 lbs	72.0	0.21	35.97	0.103
Scenario 2 45 minute ground power capacity	29.9"x19.7"x19.7"	6.72 ft ³	1235 lbs	50.6	0.15	37.77	0.111
APU Starting	15.2"x4.33"x4.33"	0.16 ft ³	31 lbs	772.0	2.3	12.87	0.041

FIGURE 6
Comparison of Chemset and F2 Plates for
Main Engine Starting Battery

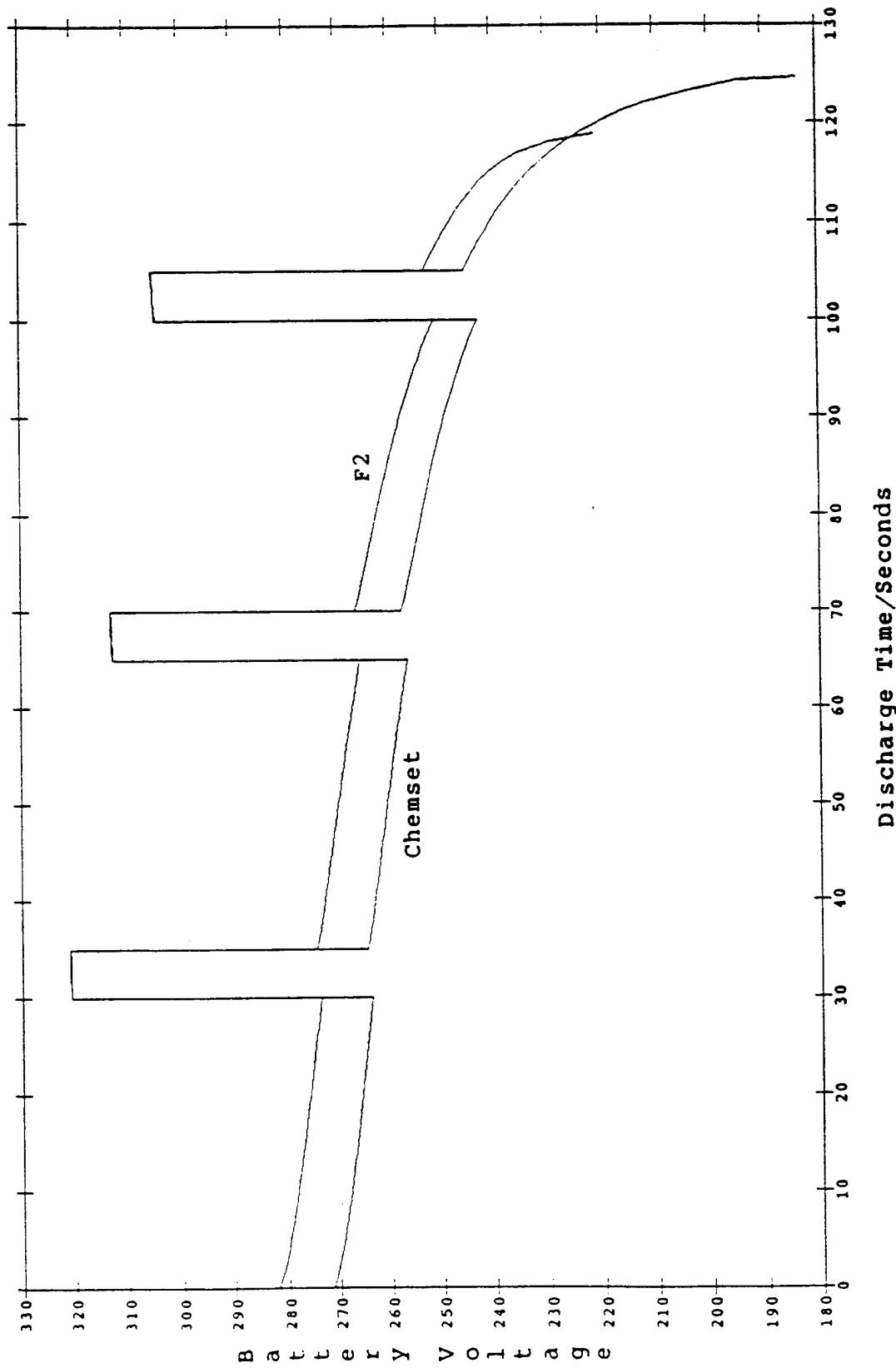


FIGURE 7
Effect of Temperature on Performance
of Main Engine Starting Battery

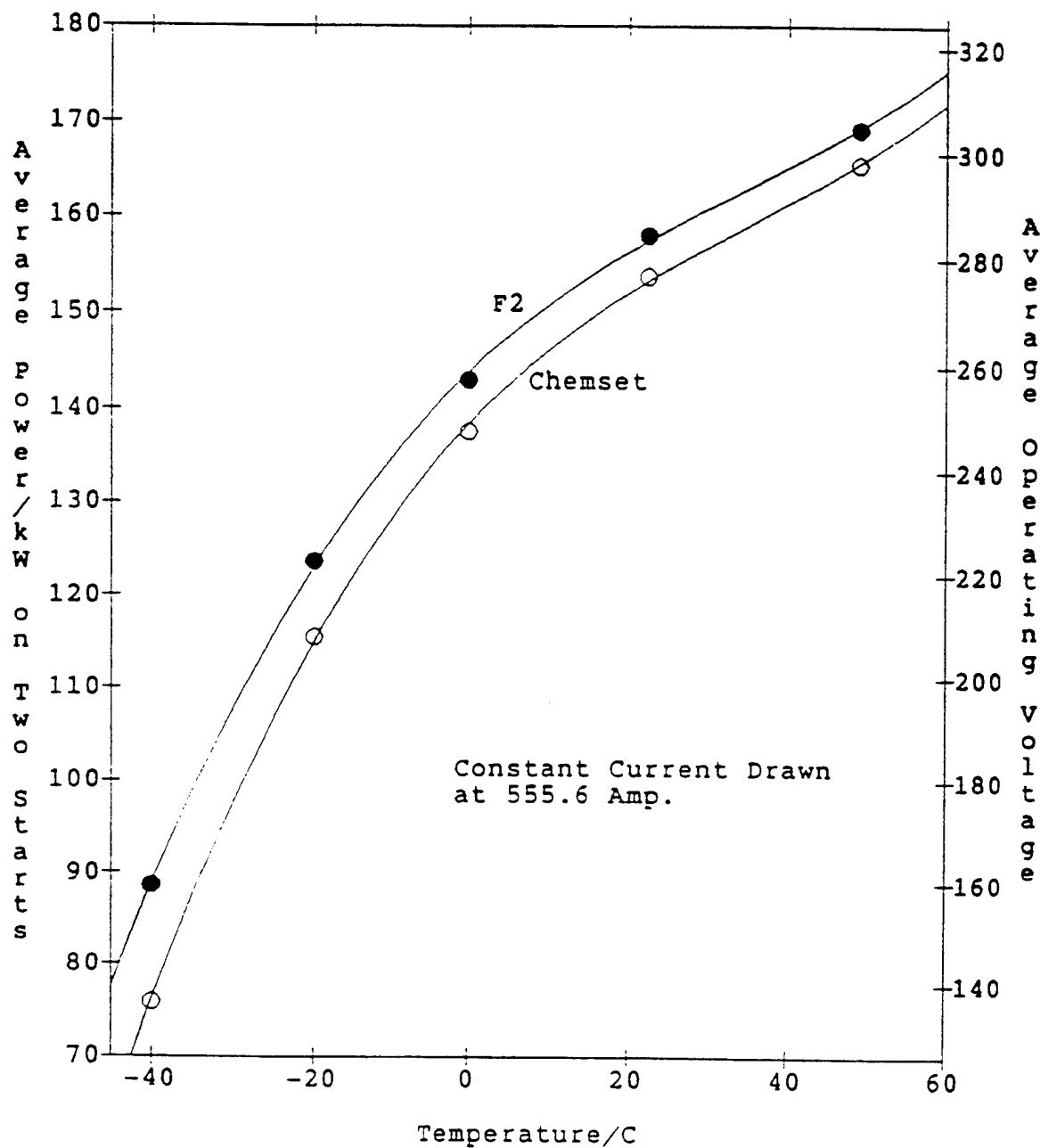


FIGURE 8
Comparison of Chemset and F2 Plates for
Ground Power Units

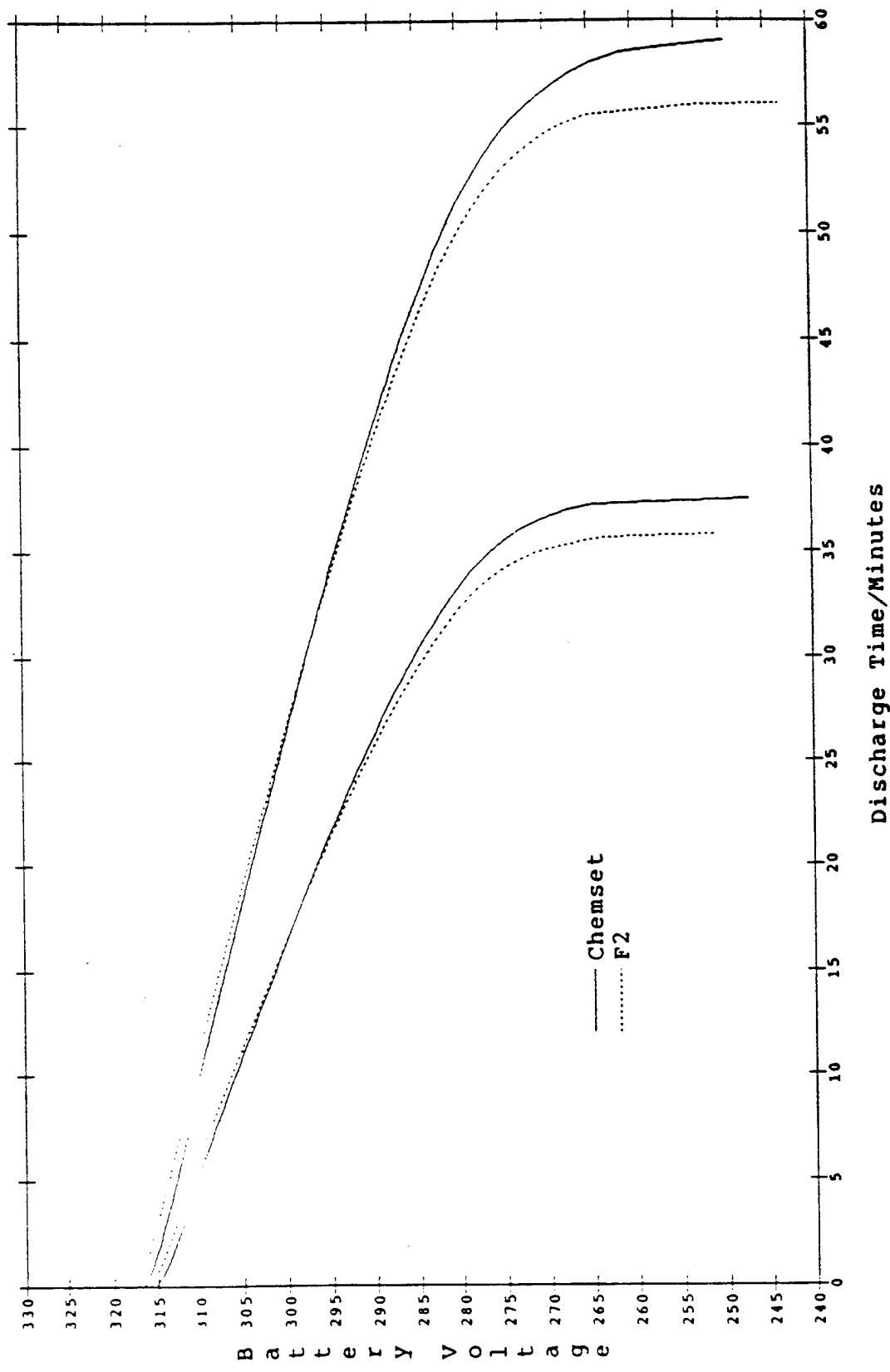


FIGURE 9
Effect of Temperature on Power Output
of the Ground Units

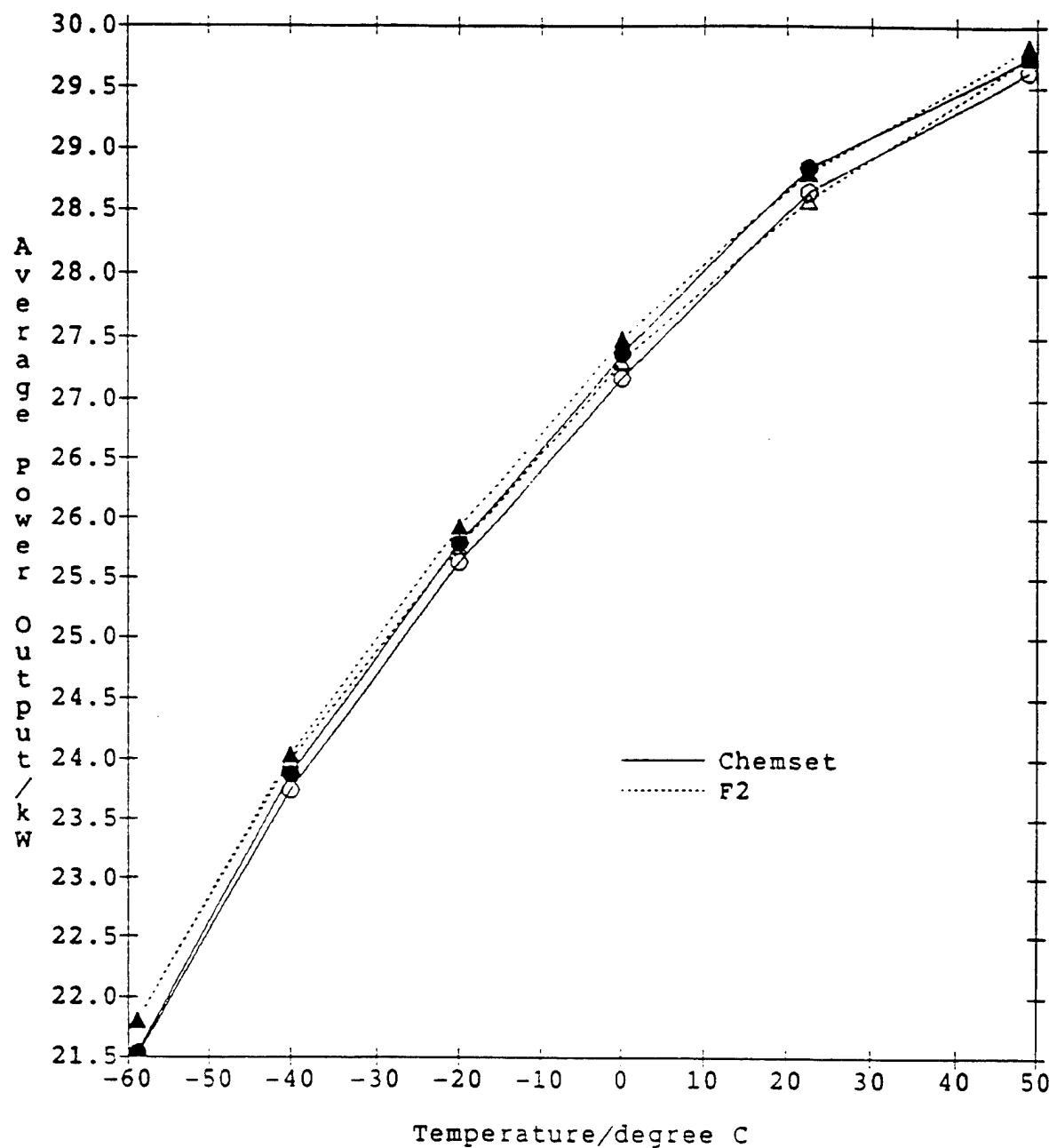


FIGURE 10
Effect of Temperature on Capacity of
the Ground Units

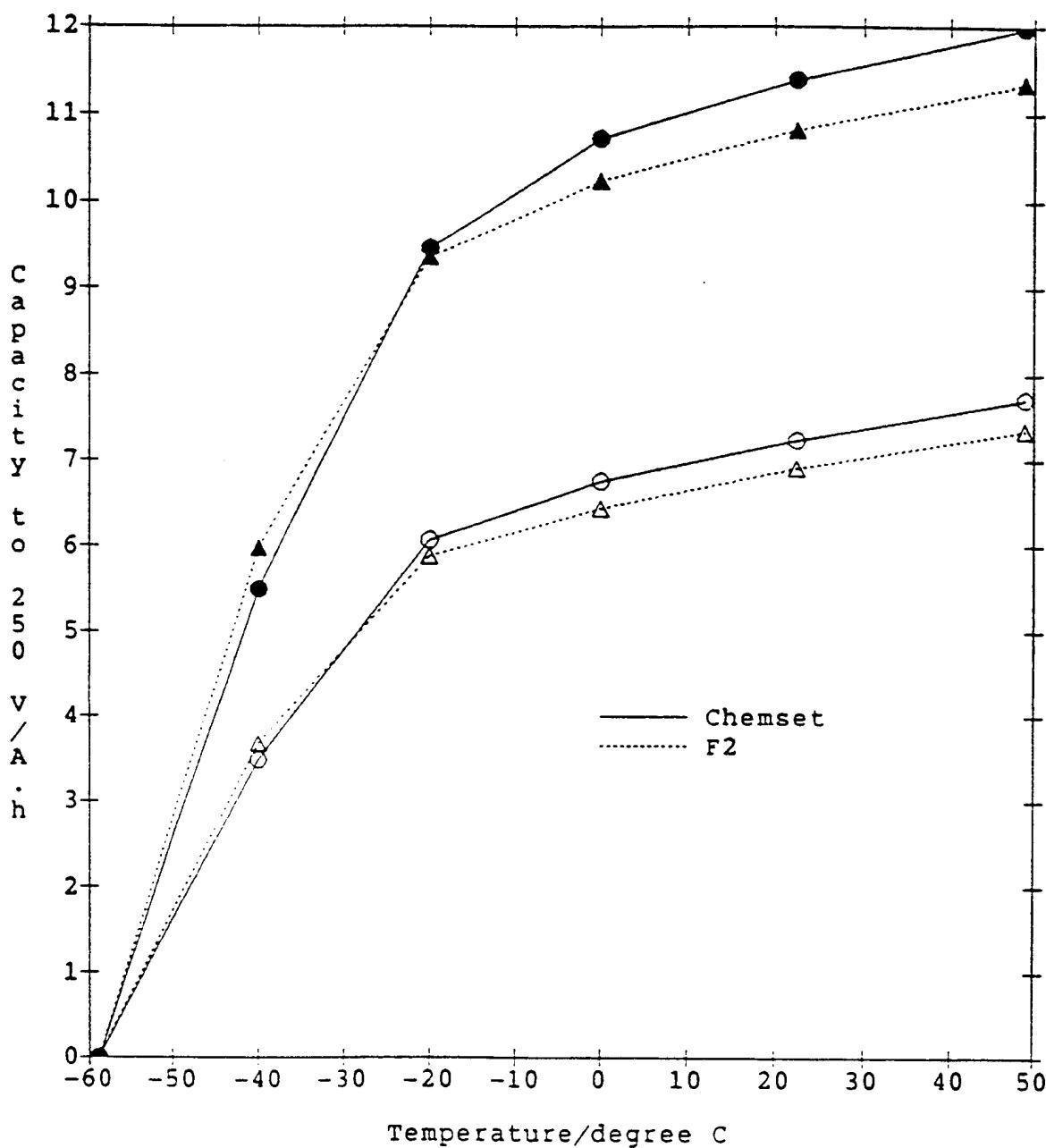


FIGURE 11
Comparison of Chemset and F2 Plates for
APU Starting Battery

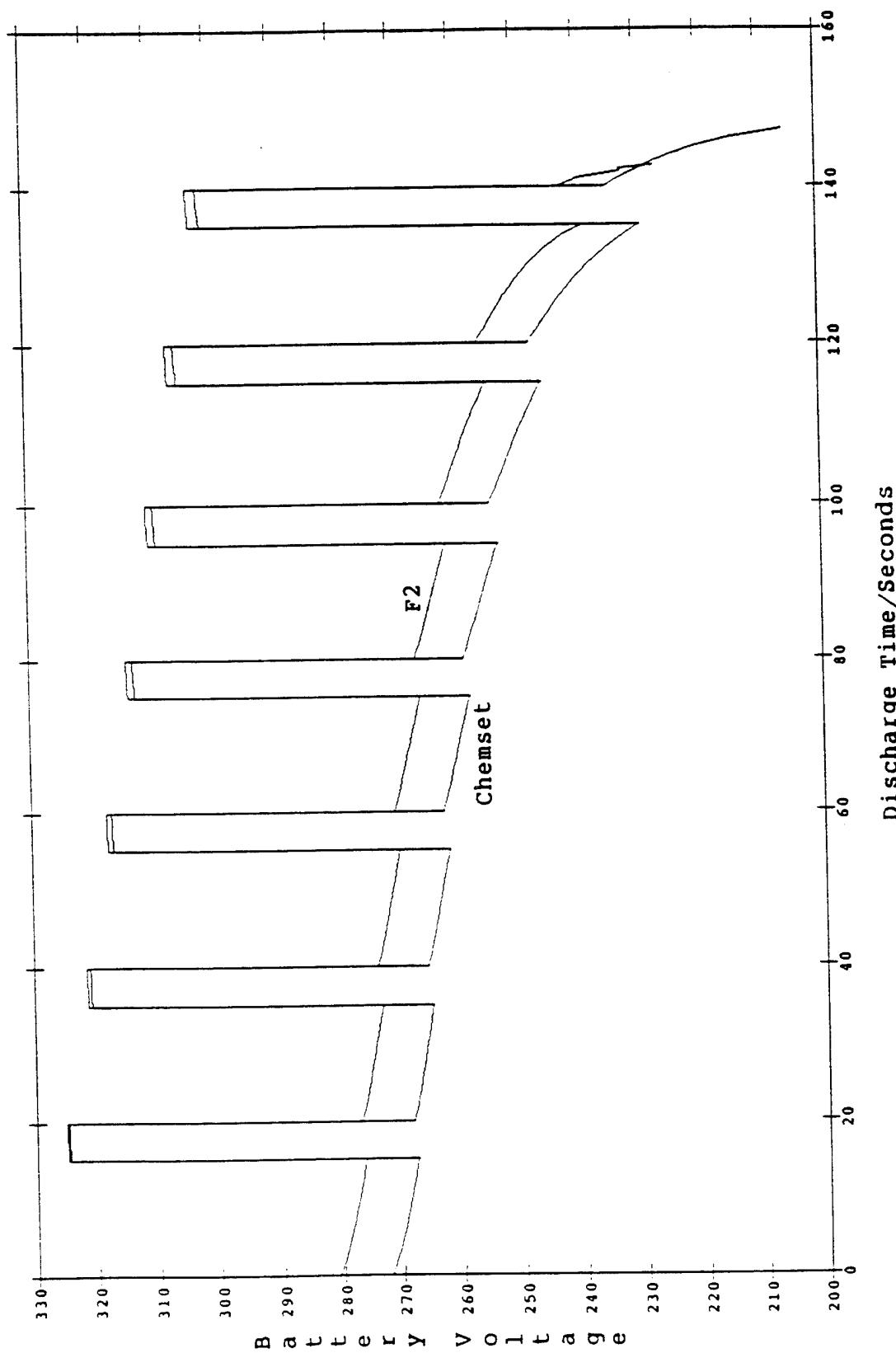


FIGURE 12
Effect of Temperature on Performance
of APU Starting Battery

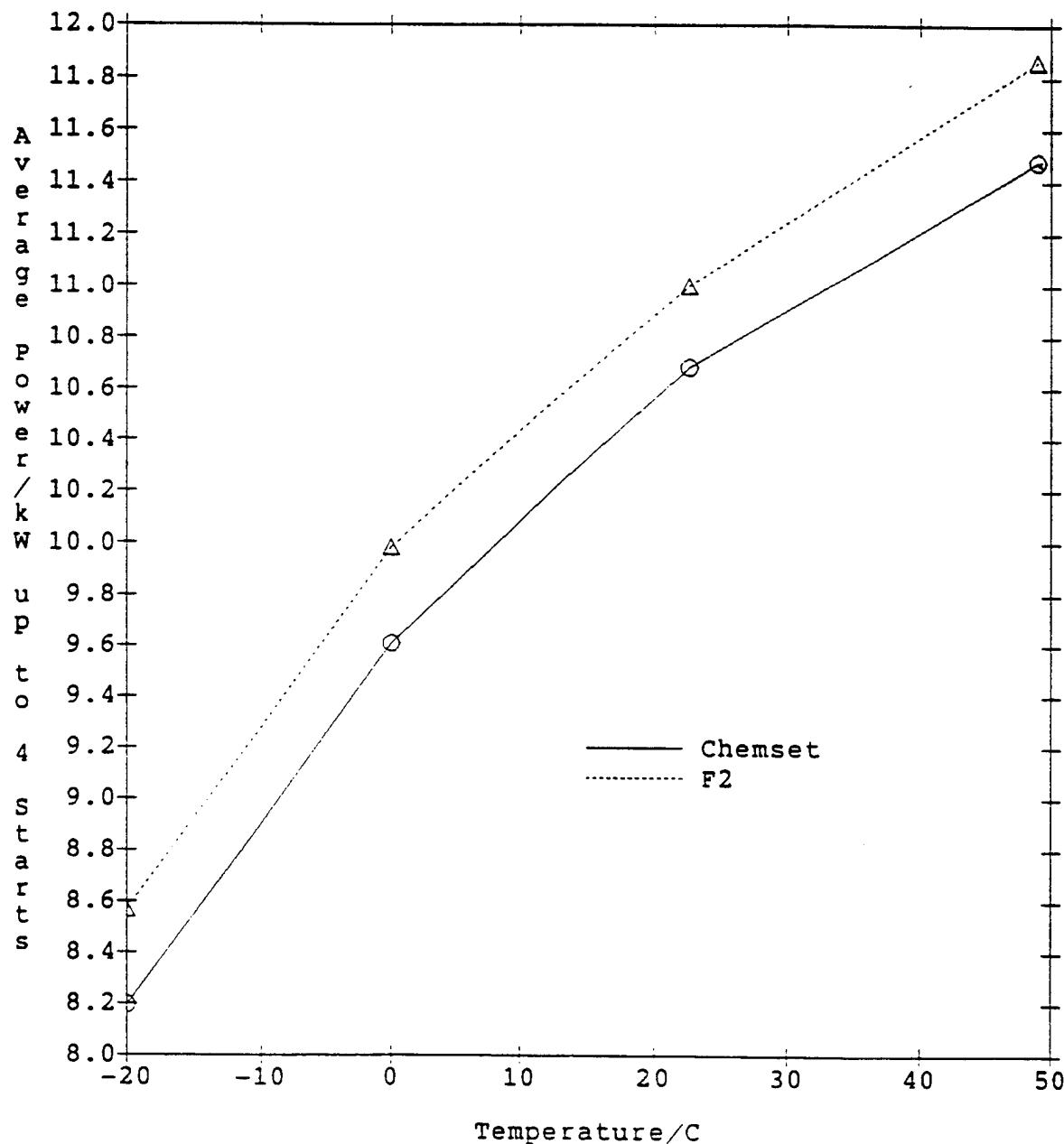


FIGURE 13
Comparison of Chemset and F2 Plates for
Emergency Power Unit

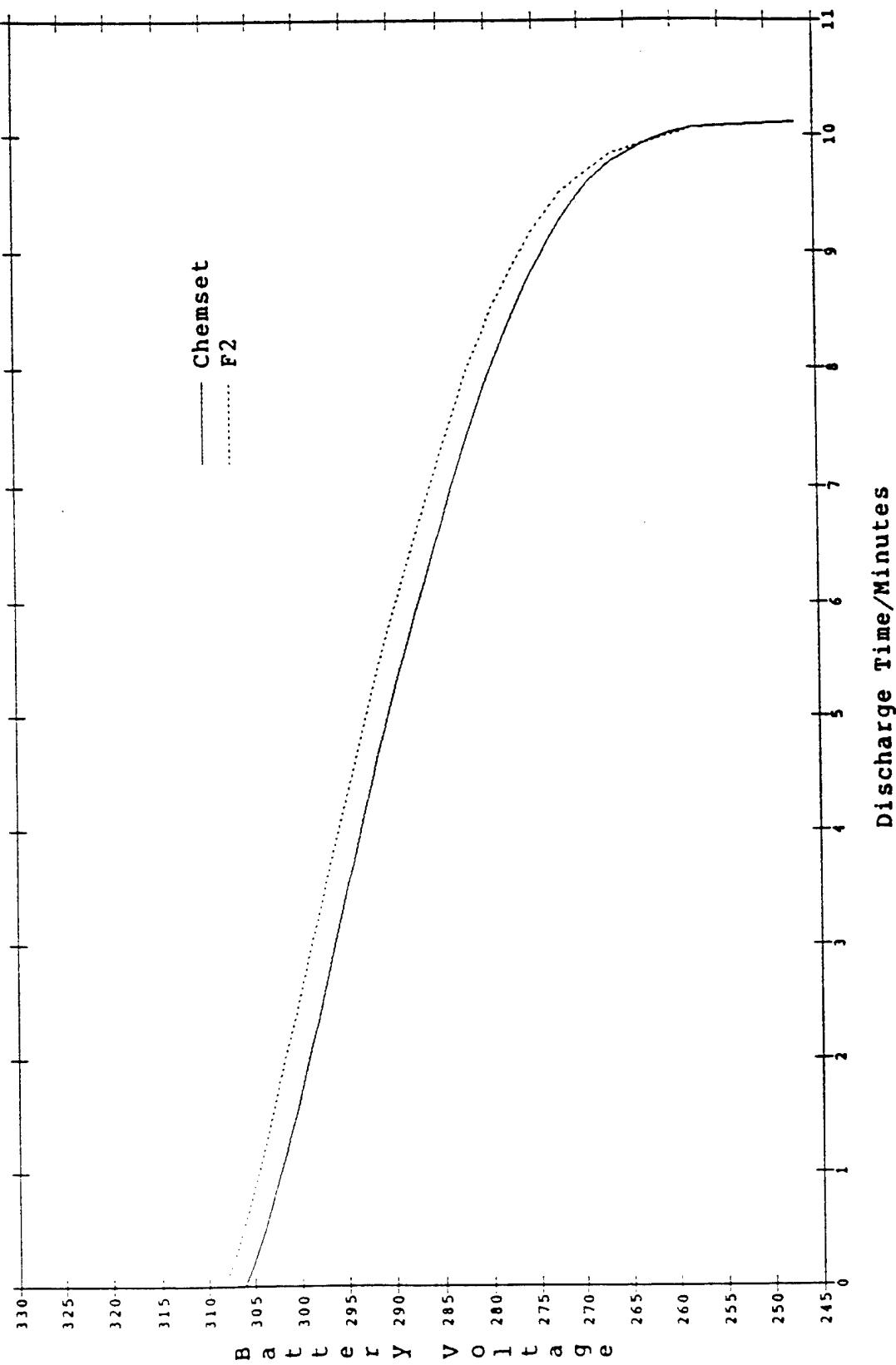
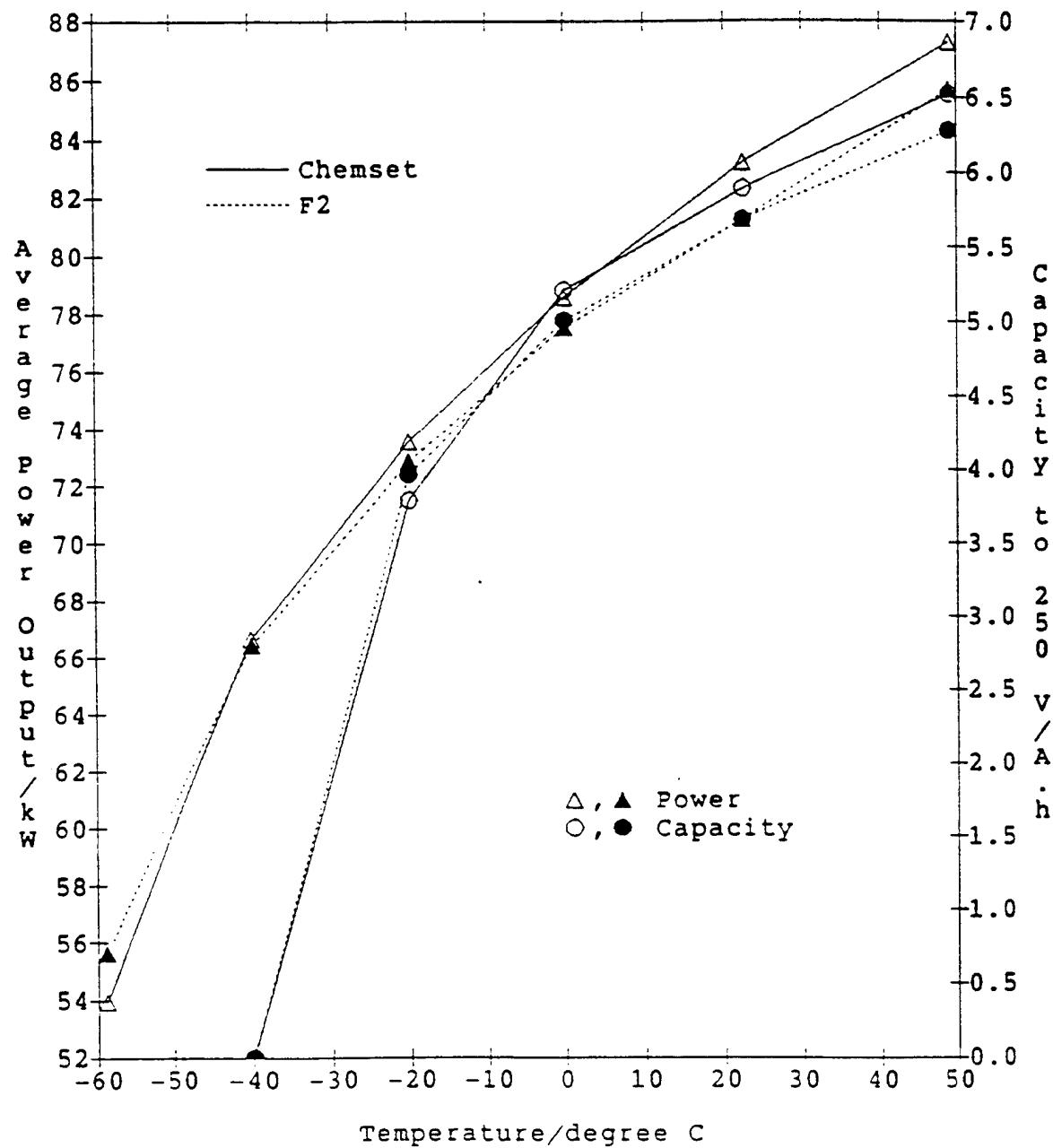


FIGURE 14
Effect of Temperature on Performance
of the Emergency Power Unit



Recognizing the recommendations from previous WPAFB work performed at JCBGI, conductive filler development resumed with further investigation of doped oxide. Coated glass fibers were also studied.

Initial work with Photon Energy Systems (PES) focused on coating doped oxide onto glass fibers. Four separate attempts were made with poor results. The first lot did not withstand the acid environment, and the second lacked uniformity and conductivity. Coated fibers from the third trial possessed no adhesion between the oxide and glass, hence were impossible to handle or compound into plastic. PES ultimately did coat 2-6" long fibers during a fourth trial, but was unable to supply the shorter lengths required for this application. Activity in this area was subsequently discontinued.

Efforts by Materials and Electrochemical Research, Inc. (MER) to produce a dense plaque of doped oxide met with similar difficulties. Prototype samples lost all conductivity and dissolved when put in contact with H_2SO_4 . A carbide compound was also provided, but found too resistive. No further attempts were made.

Two companies were next contacted for samples of doped oxide in powder form. Provided materials were extremely similar in particle size and appearance, and remained stable throughout acid leach testing. Replicate samples of 85% and 90% loaded plastic were then prepared. Measurements showed the oxide from Magnesium Elektron, Inc. (MEI) to be seven to fifteen times more conductive than that obtained from Crystal Research, Inc. (CRI). Throughout ensuing months, MEI recognized the product potential, entered into a joint development (JD) effort with JCBGI, and supplied over \$110,000 worth of oxide to JCBGI at no cost. Leftover material was returned per the appropriate clause in the JD. Additional oxides doped with other elements were prepared by MEI late in the contract, but shown highly resistive and unstable during JCBGI testing. MEI was also instrumental in providing compounding expertise that greatly expedited the development effort.

Particle size optimization was one such area in which MEI provided invaluable help. JCBGI initially believed that a smaller particle size (1 micron) would reduce porosity due to its being more easily wetted by the surrounding plastic resin. Trials using fines screened from the supplied material proved the contrary with regard to both conductivity and porosity. Resistivity readings increased twenty-fold. Discussions with MEI's compounding experts revealed that the use of uniformly shaped, ultrafine particles made it more difficult to achieve the needed particle-to-particle chain of contact through the thickness of the material, i.e. increased resistance. The smaller particle size also increased the available surface area at which pores could and did develop. All contract work was performed using particles roughly 3-5 microns in diameter. Use

of the estimated 10-20 micron optimum particle would have required an entirely different production method. Time and associated costs of the changeover were far beyond the scope of this program.

Subsequent electrical testing of the MEI material showed the doped oxide to lack stability at negative electrode potentials. This finding required doped oxide be used as a laminate in conjunction with a material better able to withstand the environment at the negative plate. Carbon black was immediately proposed as the ideal partner, having been previously identified as highly conductive, lightweight, readily available and stable at negative potentials during the first WPAFB contract. Compounding trials optimized the loading, resulting in highly conductive parts that were also very flexible.

Compounding descriptions and the corresponding conductivity measurements are provided as figures in the text.

3.3.2 Subtask 3.2 Substrate Fabrication Techniques

Given the limited batch size and trial-to-trial variability in hand compounding plastic and filler, resins were carefully chosen for study. These included low-density polyethylene (LDPE), fluoropolymer formulations (Kynar), polytetrafluoroethylene (PTFE), and high-density polyethylene (HDPE).

Given its use in prior WPAFB-sponsored work, initial efforts focused on LDPE and Microthene™ from Quantum Chemical Corporation was purchased. A powdered form was requested and received to facilitate uniform filler dispersion with minimum porosity. Dry mixing of the filler and resin was accomplished by hand using a mortar-and-pestle early on in the contract. This was later replaced by V-blending. The mixture was then melt blended in a twin screw extruder to produce pellets that were compression molded into sheet form. Early samples were thick (0.070") and used exclusively for proving the stability of the filler. After several successful resistivity tests, work was redirected on thinning the part and making it more conductive.

Another resin, PTFE, was investigated concurrently. Loadings from 70-75% produced highly conductive parts, however, these were also very porous. Investigations were undertaken with Imprex, Inc. to impregnate the porous parts under vacuum with a polycarbonate-based liquid resin to reduce the porosity without hindering the conductivity. PTFE development was stopped when samples were shown to have remained porous and become even more resistive following treatment.

Kynar was also explored for use as a base resin. The material showed initial promise, during producing conductive and nonporous material during hand compounding trials. However, the 375°C temperature needed to soften and melt the resin degraded the doped oxide. LDPE and Kynar blends resulted in conductive but highly porous material. Development in this area was discontinued given the successes with LDPE.

Additives were next employed to improve the physical properties of the substrate. Coupling agents, oils, acids, acetates and silicon compounds were each investigated in an attempt to improve part conductivity, reduce porosity, and/or improve manufacturing. Coupling agents, designed to bond the filler and surrounding base resin, offered the only quantifiable advantage. Of particular note was a coupling agent available through Kenrich Chemical, Incorporated. Additions substantially improved the resultant substrate's physical properties. Order of addition was also found critical to the end product. Greatest effectiveness was had in dry mixing with doped oxide prior to adding LDPE powdered resin.

Lastly, JCBGI investigated HDPE resin in an effort to widen the operating temperature range of the battery. Initial stability tests showed high porosity levels. Increasing the melt blend temperature produced stable parts. Development was halted in June 1994 when the program's technical direction was changed (see Section 4.0 - Metallic Substrate Development).

Alternative methods of producing sheet stock were also investigated. Molded Rubber and Plastics (MRP) and JCBGI teamed to design a vacuum compression mold to remove trapped gases and produce pore free parts. Unfortunately, samples exhibited physical properties no better than parts made in the conventional manner. Work was discontinued due to the prohibitive \$75/part cost and the large volume of material needed per trial (10+ pounds).

Skiving was no more successful. Thin rolls of doped oxide in Microthene™ were received from DeWal Industries in May 1993 for laminating and resistivity testing. Resultant laminates were 0.029-0.031" thick with resistivities in the range of 1.7-2.0 Ω-cm. Given the promise of the materials produced by DeWal's skiving process, JCBGI twice supplied additional compounded materials for processing into sheet. Doped oxide samples exhibited low initial porosities that increased as a result of the laminating process; the porosity of the carbon black material was never acceptable. Work with DeWal was subsequently discontinued.

Carbon-black development proceeded more quickly with the aid of JCBGI's zinc-bromine battery development program. Several different types of carbon-black were screened and a Ketjenblack material from Azko Chemical was chosen. Compounding trials identified an optimum carbon-black loading level that afforded parts with a conductivity of 1-1.6 ½-cm and enough flexibility to be used as a bipolar substrate.

Laminating the filled LDPE substrates was next addressed. Early laminates exhibited a resistivity higher than the sum of the constituent pieces due to the "skin" formed on the surface of each sheet when molded. Two methods of removing the "skin" were tried. The addition of carbon black at the interface prior to laminating proved effective, but difficult to perform in a uniform manner. The second and adopted method required gentle sanding of the skinned surfaces with sandpaper. Sanding prior to lamination resulted in a 50-75% reduction in part resistivity and no effect on part stability.

3.3.3 Subtask 3.3 Stability Testing

The procedure and fixture for quantifying a bipolar substrate's stability in acid and under potential were developed at JCBGI over many years. Both three- and four-point tests were required to evaluate a sample's viability.

As shown in Figure 15, a substrate sample was clamped between two hollowed polycarbonate endblocks, exposed to electrolyte, and wired as the working electrode. A potential of 1.5 volts was applied and the current collected at the top of the substrate in the three-point system. After 24 hours on test to establish a baseline current, the leads were rearranged to collect current after passing through the substrate, i.e., the four-point test. The test continued for a minimum of 3 additional days. No change in the current acceptance established the sample to be nonporous. A rising current suggested porosity or filler instability. Detailed stability results are provided in Appendix B.

Conductivity before and after the three- and four-point regimen was also monitored. An increase of 20% or more signalled porosity or filler instability. Since doped oxide had been successfully tested, an increase in resistivity was interpreted as increasing porosity, i.e., carbon-black was exposed to the positive potential as a result of the porosity causing the carbon-black to oxidize and become nonconductive.

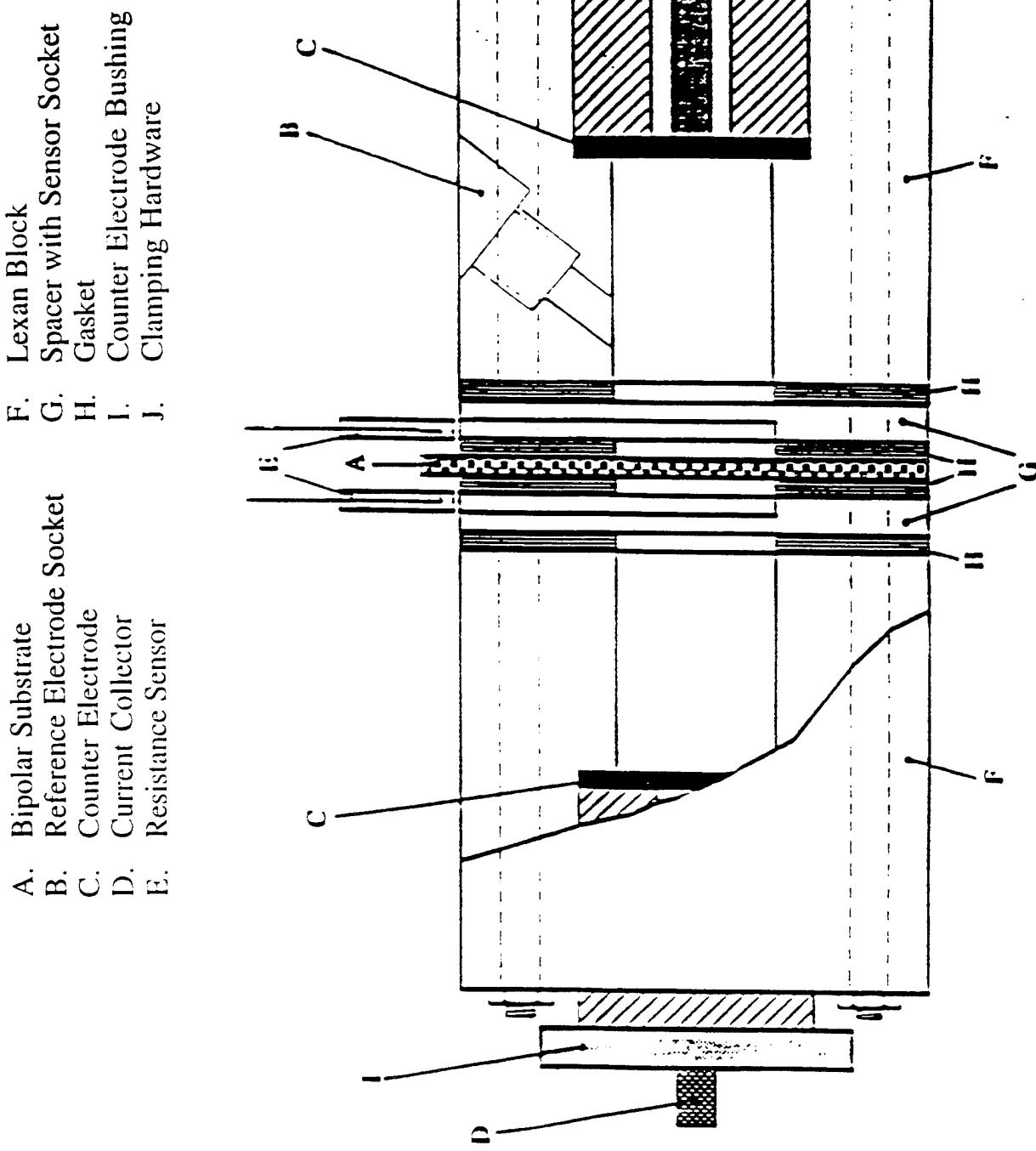
3.3.4 Subtask 3.4 Proof of Concept Testing

Over 60 batteries of various voltages were assembled and tested. Dry, unformed electrodes with 10 in² active areas were alternately stacked with elastomeric spacers and compressed to dimension between 0.5" thick polycarbonate end plates. Insulated bolts positioned around the perimeter of the fixture were easily tightened to compress the gaskets to affect hermetic cell seals. Absorptive glass mat separator was placed between opposing electrodes and filled with electrolyte through channels machined across the upper portion of each

FIGURE 15

Stability Test Fixture

- A. Bipolar Substrate
- B. Reference Electrode Socket
- C. Counter Electrode
- D. Current Collector
- E. Resistance Sensor



gasket. Discharge performance routinely surpassed 5 minutes at 1 A/in², but with limited cycle life.

Laminate and positive paste adhesion were the ultimate issues and numerous approaches were investigated in attempts to foster them. Techniques included roughening the pasted surface with various grit sandpaper, embedding fibers, sintering lead dust or oxide powder onto the active areas, flame spraying lead, pretreating the plastic to increase its wettability. A review of the battery build sequence, documented in Figure 16, quickly shows that any battery formed without the use of lead sheet could not be tested due to high internal resistances caused by poor paste adhesion.

The major breakthrough occurred upon recognizing the special needs of polyolefins. Involved surface pretreatments are recognized as necessary to achieve bonds with wax-like surfaces that are difficult to wet if left alone. Surface treating LDPE prior to attaching a layer of thin lead foil decreased the part resistivity by 50-75%. Over 150 cycles were demonstrated with shorting as the cause of failure. Subsequent builds neared this benchmark, however, lead foil delamination became a recurring problem. Substrate conductivities checked prior to pasting and after cycling showing no change added to the confusion. Treatment parameters were reviewed and found incorrect, resulting in delamination *within* the plastic part. Optimization trials were initiated, along with investigations of HDPE resin. HDPE was proven to bond more strongly to lead sheet, but the resulting cycle life was still unacceptable. Efforts were halted with the change in the program's technical direction.

3.4 WBS 5.0 BATTERY FABRICATION

3.4.1 Subtask 5.1 Sealing Methods

Two 10-volt batteries were produced using an injection molded containment method in October 1993. Electrodes, separators and spacer frames were arranged to form a stack that was inserted into a cavity for molding. Plastic injected into the mold formed a frame around the entire stack to provide the necessary sealing and spacing requirements, as well as provisions for acid fill.

Electrode quality within each 10-volt stack was poor due to the required part size. Length and width exceeded the working area of the press. Pieces were 0.080" thick and highly resistive (10 Ω -cm). Cross sectioning of one dry, unformed (DUF) stack showed complete plastic fill and no electrode distortion. Confirmation of hermetic cell-to-cell sealing was never

FIGURE 16
Composite Battery Builds

<u>ID</u>	<u>Volts</u>	<u>Adhesion Method</u>	<u>Cycles</u>	<u>Cause of Failure</u>
159	4	Lead dust	32	Lack of paste adhesion
159-B	4	Lead dust	15	Lack of paste adhesion
160	4	Lead dust	15	Lack of paste adhesion
182-1	4	Lead dust	5	PbSO ₄ at surface
182-2	4	Sanded surface	5	Lack of paste adhesion
182-3	4	Lead dust	5	PbSO ₄ at surface
182-4	4	Sanded surface	5	Lack of paste adhesion
194-3A	4	Embedded 0.003" glass mat	0	PbSO ₄ at surface
194-4A	4	Embedded 0.003" glass mat	0	PbSO ₄ at surface
194-3A	4	Finely sanded surface	0	PbSO ₄ at surface
194-4A	4	Finely sanded surface	0	PbSO ₄ at surface
205-1	4	0.001" perforated lead foil	18	Lack of paste adhesion
205-2	4	0.001" perforated lead foil	18	Lack of paste adhesion
205-3	4	0.001" lead foil	19	Lack of paste adhesion
205-4	4	0.001" lead foil	14	Lack of paste adhesion
214-1	4	0.010" lead foil over treated surface	18	Leak, cracked substrate
214-4	4	0.010" lead foil	21	Lead foil delamination
214-5	4	0.010" lead foil	45	One very dry cell
214-6V	6	0.010" lead foil over treated surface	47	Lead foil delamination
218-1	4	Carbide fibers	2	Too resistive to cycle
218-2	4	Carbide fibers	2	Too resistive to cycle
224-4	4	0.010" lead foil over treated surface	151	Shed PAM, shorting
224-5	4	0.010" lead foil over treated surface	104	Lead foil delamination
241-2	4	Flame sprayed lead	0	High IR, no AM adhesion
242	12	0.005" lead foil over treated surface	15	Lead foil delamination
242-4	4	Paste over treated surface	0	High IR, no AM adhesion
243-6V	6	0.005" lead foil over treated surface	12	Lead foil delamination
257	12	0.005" lead foil over treated surface	8	Lead foil delamination
259	12	0.005" lead foil over treated surface	0	Lead foil delamination
260-2	4	0.005" lead foil over treated surface	19	Lead foil delamination
263	6	0.005" lead foil over treated surface	9	Lead foil delamination
265	6	0.005" lead foil over treated surface	4	Crack, leak, delamination
267-1C	4	0.005" lead foil over treated surface	15	Lead foil delamination
267-4P	4	0.005" lead foil over treated surface	135	Local lead foil delamination
267-5P		0.005" lead foil over treated surface	13	Local lead foil delamination
267-6VP	6	0.005" lead foil over treated surface	33	Local lead foil delamination
267-6P		0.005" lead foil over treated surface	18	Local lead foil delamination
267-8C	4	0.005" lead foil over treated surface	20	Local lead foil delamination
267-9P	4	0.005" lead foil over treated surface	20	Local lead foil delamination
267-11C	6	0.005" lead foil over treated surface	11	Local lead foil delamination
268-6VC	6	0.005" lead foil over treated surface	11	Local lead foil delamination
268-8C	4	0.005" lead foil over treated surface	9	Local lead foil delamination
268-10C	4	0.005" lead foil over treated surface	68	Local lead foil delamination
268-11C	4	0.005" lead foil over treated surface	135	Local lead foil delamination
268-12C	12	0.005" lead foil over treated surface	15	Lead foil delamination
277-1C	4	0.005" lead, treated surface, acid dip	2	Local lead foil delamination
277-2C	4	0.005" lead, treated surface, acid dip	4	Local lead foil delamination
277-6VC	6	0.005" lead, treated surface, acid dip	3	Local lead foil delamination
278-1C	4	0.005" lead, treated surface, acid dip	8	Local lead foil delamination
281-1	4	0.005" lead on HDPE, treated surface	6	Local lead foil delamination
282-1	4	0.005" lead on HDPE, sanded, treated surface	23	Lead foil delamination
282-2	4	0.005" lead on HDPE, sanded, treated surface	5	Lead foil delamination
282-6V	6	0.005" lead on HDPE, sanded, treated surface	5	Lead foil delamination
285-1	4	0.005" lead, washed oxide, treated surface	11	Lead foil delamination
286-2	4	0.005" lead, unwashed oxide	0	Short
286-3	4	0.005" lead, unwashed oxide	11	Lead foil delamination
287-2	4	0.005" lead on HDPE, washed, treated surface	1	Cracked substrate
287-3	4	0.005" lead on HDPE, treated surface	10	Lead foil delamination
287-4	4	0.005" lead on HDPE, treated surface	6	Cracked substrate

obtained due to difficulties porting the cells for pressurization tests. The trial did, however, prove that injection molded containment was a viable manufacturing technique.

4.0. METALLIC SUBSTRATE DEVELOPMENT

4.1 WBS 1.0 PROGRAM MANAGEMENT

4.1.1 Subtask 1.1 Managing Strategy

Effective July 28, 1994, Ms. Jennifer Rose assumed the responsibilities of the contract's previous project engineer, Mr. Doug Pierce, due to his departure from JCBGI.

Shortly thereafter, a proposal requesting a no-cost time extension was submitted to the contract negotiator on July 13, 1994. Gantt charts detailing this effort are shown in Figures 17 and 18. This followed a discussion with Mr. Richard Marsh during which it was mutually agreed that, despite significant advances in composite bipolar substrate development, remaining WPAFB contract work should be focussed on the use of a lead substrate with improved corrosion resistance. Through a parallel bipolar program, JCBGI had repeatedly demonstrated 2000+ cycles in a 12-volt configuration utilizing lead substrates, and over 5700 cycles using a 6-volt unit. Laminated metallic substrate work had also been underway for nearly 12 months in an effort to increase corrosion resistance, and hence, cycle life.

4.2 WBS 2.0 BATTERY DESIGN

4.2.1 Subtask 2.1 Battery System Design Analysis

The existing small metallic bipolar battery design was scaled up and modeled to investigate high power performance. Results suggested the use of a thinner cell design to be critical to achieving rates of 500 W/kg and higher. Per these findings, work was redirected to designing a 24-volt module within the volume previously allotted for 12-volts. This effectively aligned the contract deliverable voltage with WPAFB's ultimate application and JCBGI's commercial product target. Constant power performance projections are shown in Figure 19.

4.3 WBS 3.0 BIPOLAR PLATE

4.3.1 Subtask 3.1 Multialloy Substrate Development

Under separate contract, JCBGI began investigations into laminated metal substrates in November 1993. Corrosion testing of three, four and five layer samples and constituent alloys was performed in a bipolar configuration to assess time to breakthrough. Unpasted samples were mounted in the previously described stability test fixtures (Composite Substrate Work, Subtask 3.3) for three-point testing. Only the positive surface was exposed to electrolyte. Working and reference electrodes were also introduced. Initial testing of a new material was

FIGURE 17: No-Cost Time Extension Gantt Chart with Milestones

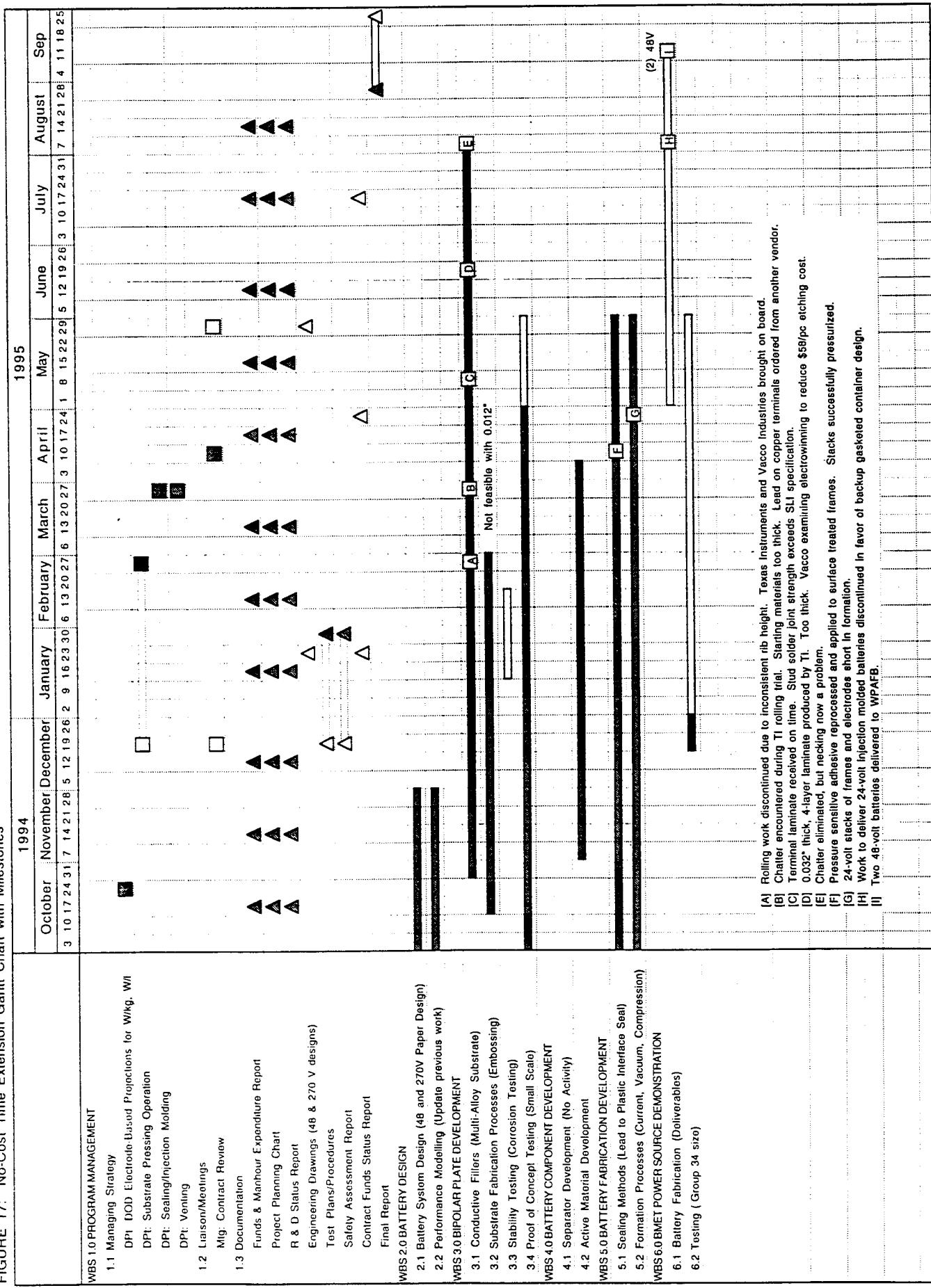


FIGURE 18: WPAFB Bipolar Deliverable Schedule

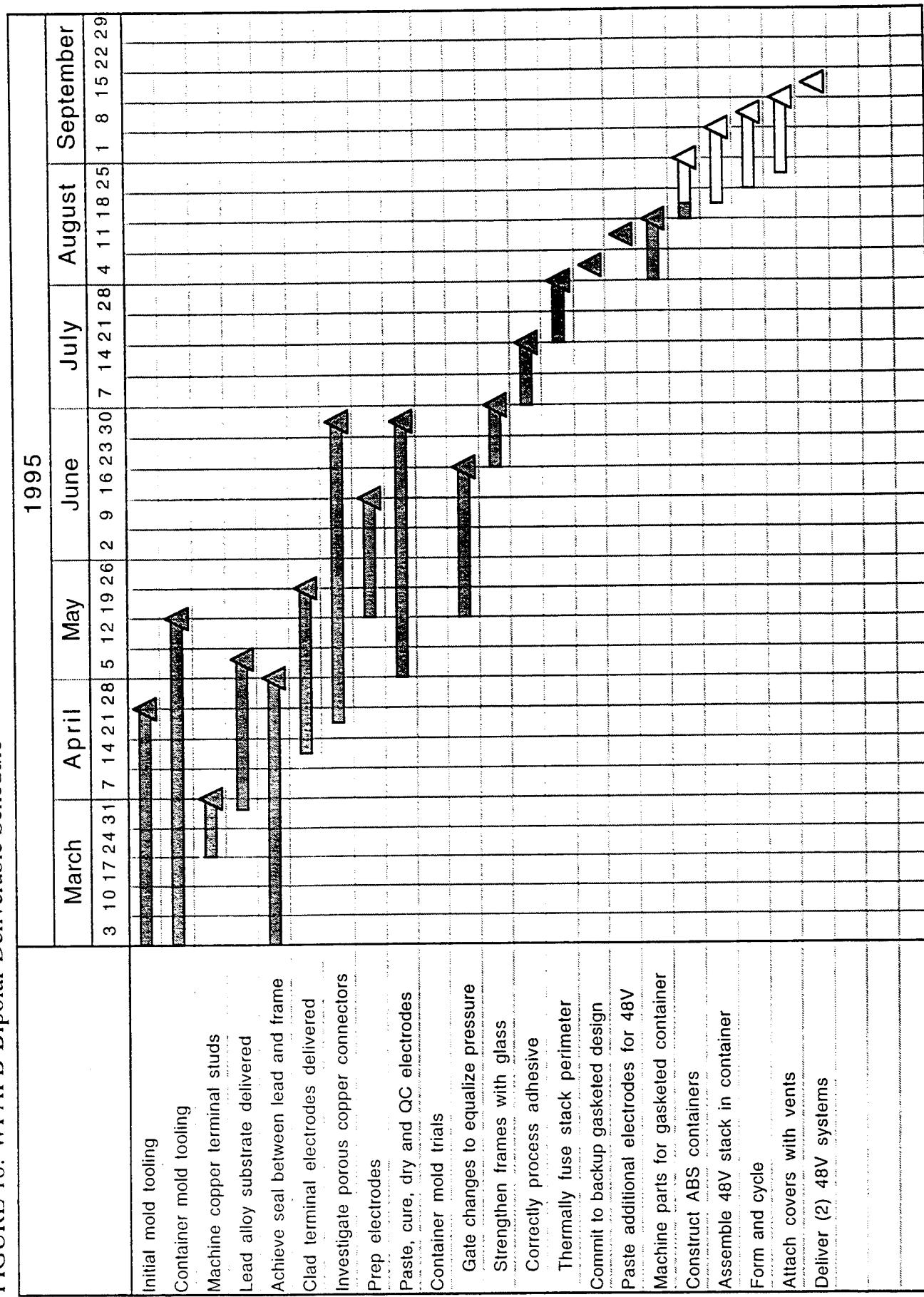
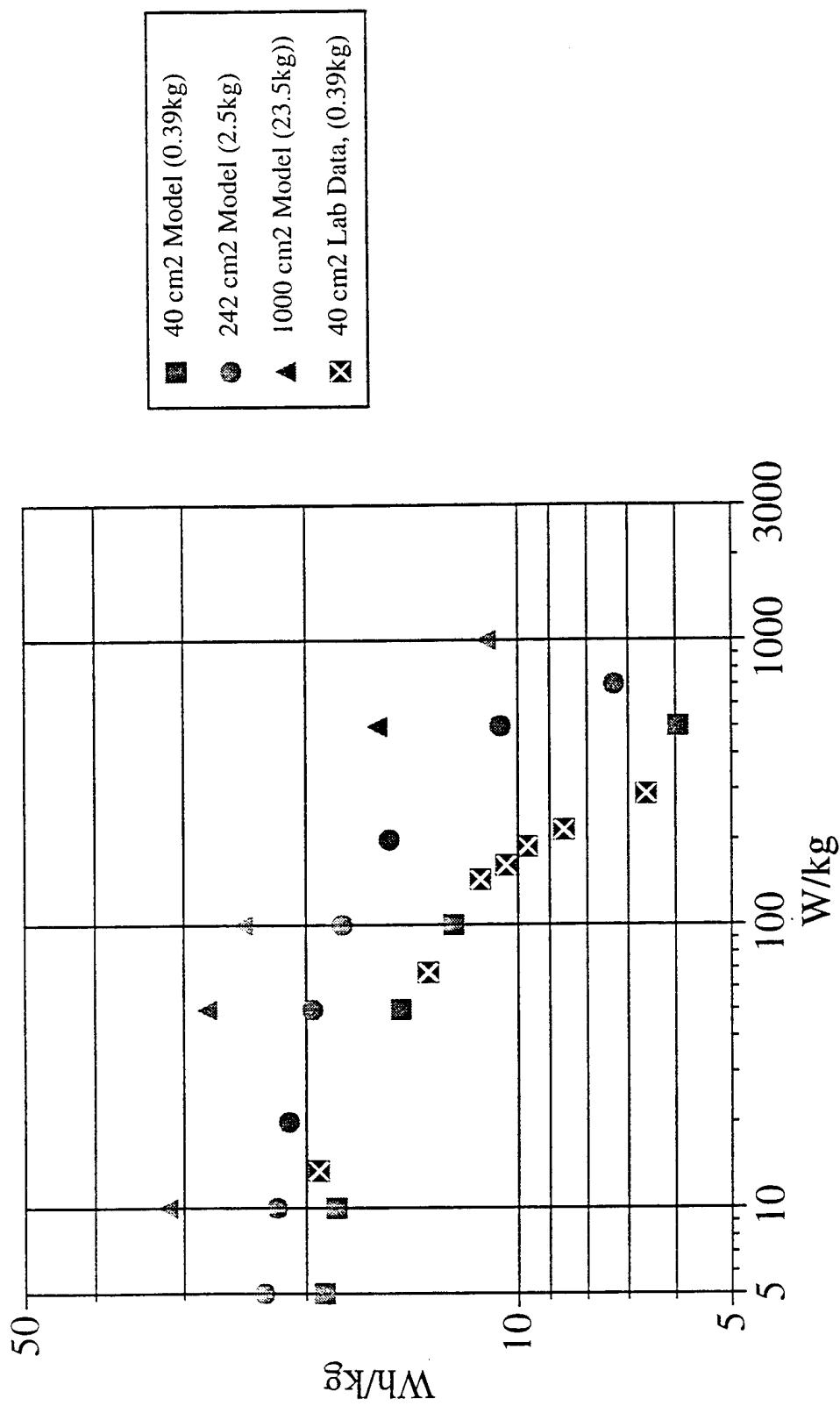


FIGURE 19
Constant Power Performance Projections
Metallic Bipolar Substrate



performed at 70°C and a constant potential of 1.50 V until evidence of pinholes was noted, i.e., liquid in the back chamber or spikes on the current acceptance curve. Replicate samples were then run, pulled at points prior to breakthrough, and submitted for cross sectional photomicrographs to quantify the corrosion rate.

Comparing rates of all samples tested showed the corrosion resistance of laminates to be second only to that of a high silver content alloy. Batteries utilizing the clad material were assembled and tested, but performance was poor. Teardowns showed improper cleaning of the starting materials to have prevented bonding of the dissimilar metals at the molecular level. Delamination resulted in high internal resistance that impeded high rate performance.

In October 1994, assistance was sought from Texas Instruments' Cladding Division (TICD), a leader in the laminating industry. Partnership activities were slow to materialize due to reorganization within TICD, however, two- and three-layer trials cladding lead to a stainless steel core were successful in December 1994. In March 1995, lead clad copper material was received and forwarded to Vacco Industries (see Metallic Substrate Work, Subtask 3.2) for surface etching trials. TI had planned bonding and rolling to facilitate a 65% reduction of the 0.054" thick constituent layers, however, a maximum of 51% was achieved before "chattering" (rippling) was observed. Secondary rolling ruined the bonds achieved in the first pass. New starting materials were requested for the production of 0.013" thick material, but the May delivery date made it unlikely that the laminated material would be available for use as the bipolar substrate in the required deliverables. Four layered, 0.032" thick sample material was received in June, and required reducing the copper core thickness by 50%. The likelihood of having the concept ready for deliverable use then dismissed.

4.3.2 Subtask 3.2 Rolling/Embossing Work

Fostering paste adhesion to metal sheet requires the surface to be roughened in some manner. Small-scale metallic substrates possessed exemplary adhesion when hot pressed in a mold to create ribs protruding from each face. The raised pattern successfully broke up the "single paste pellet" that would otherwise sheet off the lead substrate during handling, and increased the surface area biting into the active material.

Substrate production times were slow and scale up required the use of more tonnage than available on any in-house press. It also lacked promise as a high speed, manufacturing process. A roller die was ordered and five hundred pounds of 0.020", 0.025" and 0.030" thick lead were delivered to MP Metal Products for rolling trials. Without authorization, MP turned to blanking the electrodes from a compression die when the first rolling trial was unsuccessful. Rolled

samples were never provided to JCBGI for evaluation. When informed of the new production direction, JCBGI reiterated their interest in the rolled concept, but conceded to whatever parts could be produced. Time was short. MP continued their effort to produce parts, but quickly found their press tonnage insufficient. Hence, a new vendor was located. Walking 300 tons force across the die produced acceptable parts from 0.020" thick starting material. Efforts to reduce the substrate thickness to the required 0.012" thickness were unsuccessful and the embossing effort abandoned.

Photochemical etching was investigated in conjunction with laminating activities (Metallic Development Work: Subtask 3.3.1). Early trials produced copper pieces that were electroplated with lead, pasted and shown to possess good adhesion. Solid lead sheet was not etched as easily, requiring strong chemicals that made the technique cost prohibitive (\$58/piece).

As backup, plastic screen was used. Pieces were cut to the size of the active material area, pressed to eliminate elevated nodes that could cause shorting through the separator, and were tacked to the lead substrate. This alternative eliminated roughly 240 grams of lead rib mass per battery, but required significantly more labor input than the embossing concept. Despite its facilitating acceptable results, the use of plastic screen is not recommended for manufacturing.

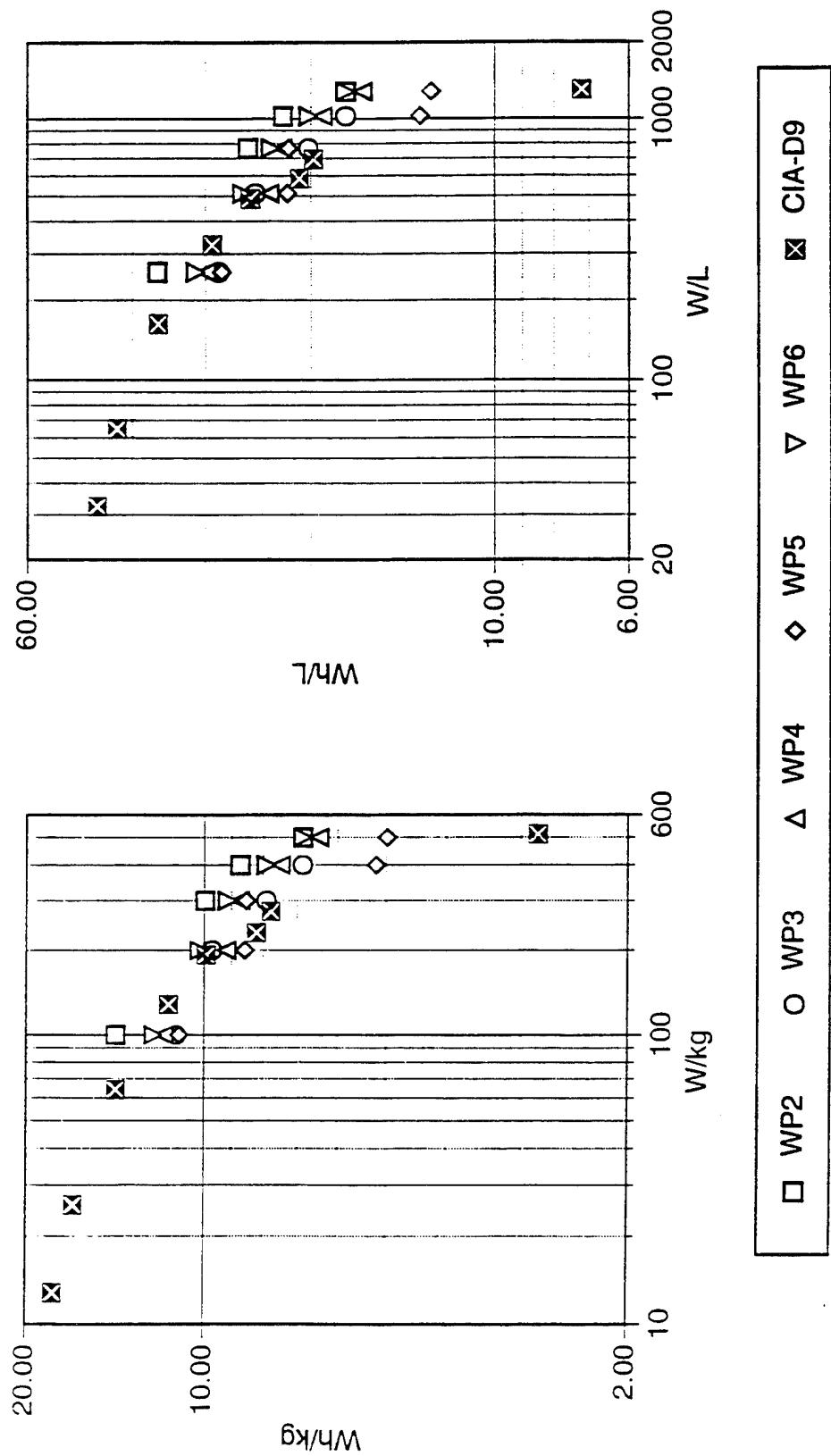
4.3.3 Subtask 3.3 Substrate Corrosion Testing

Laminates received from Texas Instruments were never corrosion tested due to their being too thick.

4.3.4 Subtask 3.4 Small Scale Characterization

Bipolar batteries having 0.012" thick substrates and 0.030" thick pasted layers were assembled, formed and tested in January 1995. Constant power discharge performance plots normalized to battery mass and volume are shown in Figure 20. Performance by WP2 and WP6 represented the best of the lot and greatly exceeded that reported for batteries delivered under the parallel metallic bipolar development contract. This was attributed to the use of 1.265 sg fill/form electrolyte. Reproducibility was an issue and investigated. Teardowns showed sulfated positives and dull negatives. Cured paste analyses reported consistently high levels of free lead that could cause initially poor or rapidly declining performance. A review of pasting procedures showed the starting PbO to be within specification and the paste code to be adequately sulfated

FIGURE 20
Constant Power Performance Normalized to Mass and Volume



and consistent from mix to mix. The dry bulb within the curing chamber was found cracked and was repaired prior to further assembly operations.

Testing of four newly-formed 12-volt units showed 10-15 cycles at 100 W/kg to be necessary to reach full capacity. Discharge times were tightly grouped after formation (Figure 21). WP-12 lagged due to oxygen ingress at cycle 3. A cursory investigation of constant current rates (Figure 22) was performed to give insight into the constant power rates required per the test plan. Constant power performance was plotted along with the modeling prediction in Figure 23, then translated into the time versus power curve shown in Figure 24.

4.4 WBS 4.0 BATTERY COMPONENTS

4.4.1 Subtask 4.2 Active Material Development

Procedures and equipment were reviewed when the free lead content in positive and negative cured plates was reported at 5.5 and 10%, respectively - far above the 4% maximum. Increasing the curing residence time from 16 to 40 hours had little effect. Moisture content was found low (6-7%) as referenced to industry and company standards and, subsequently, paste code and plate handling techniques were reviewed. Efforts to keep plates moist while awaiting transport to the curing chamber only slowed the cure reactions and actually increased the cured free lead content. Lastly, the ABR humidity chamber was diagnosed with a cracked dry bulb, repaired and reset. Cured positive and negative plates from eight subsequent pasting runs displayed acceptable free lead content following a 24 hour residence time in the environmental chamber.

A limited investigation into the effects of freezing and thawing a small 12-volt battery was performed. One unit was tested at room temperature to establish a baseline capacity and then chilled to -60°C. A 5-hour thaw was allowed and the discharge test repeated. Evidence of cell reversal and a 13% capacity loss was documented. Confirmatory work was placed on hold to allow pasting, stacking and debugging of the formation techniques proposed for full-size, 24-volt units.

4.5 WBS 5.0 BATTERY FABRICATION

4.5.1 Subtask 5.1 Sealing Methods

A variety of compounds was evaluated for use in achieving a hermetic cell-to-cell seal. In the end, an engineering sample of hot melt adhesive was pressed between release paper into

FIGURE 21
Small Scale Characterization
Capacity Development, 24 deg C

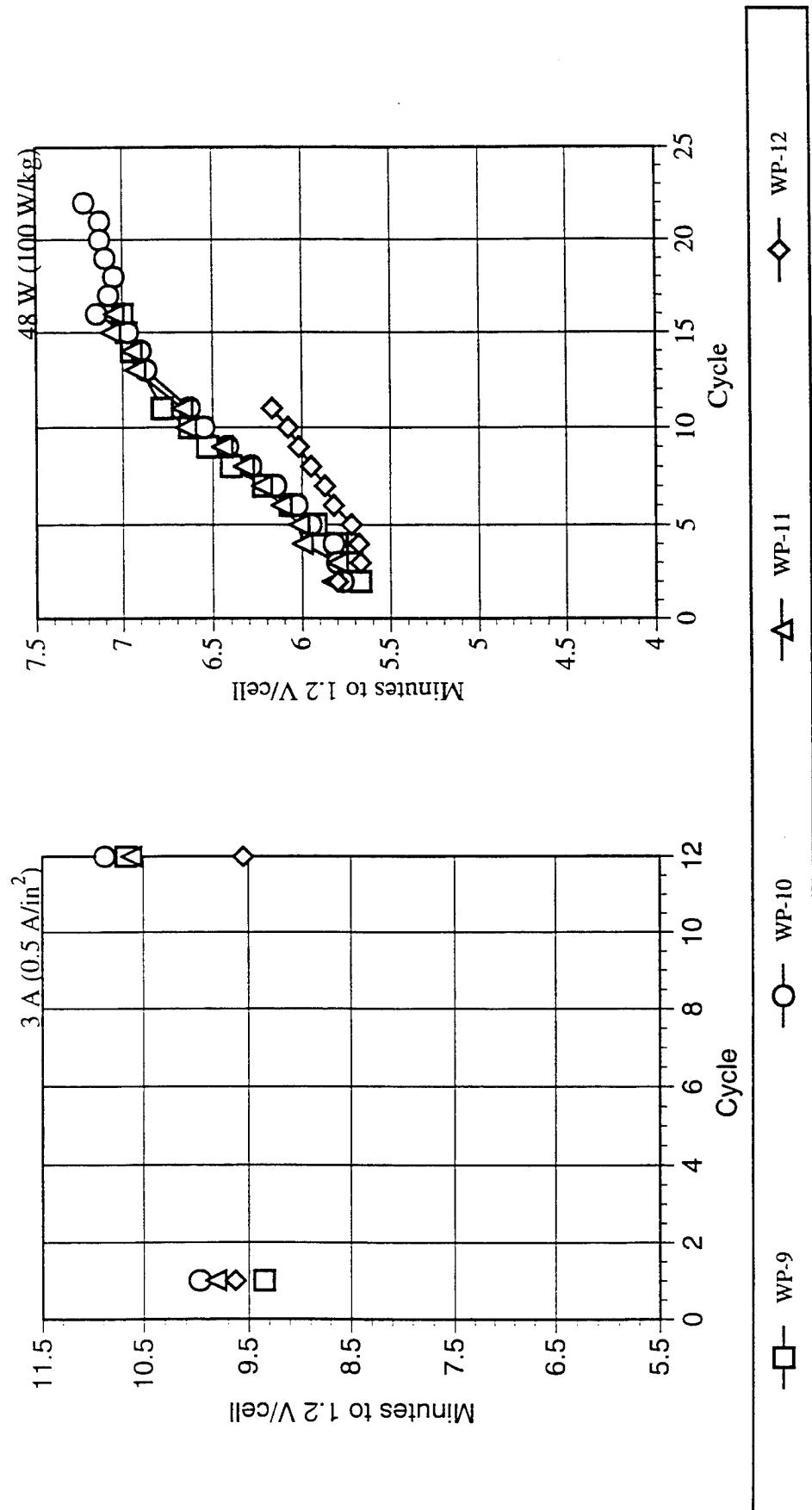


FIGURE 22
Small Scale Characterization
Peukert Relationship, 24 deg C

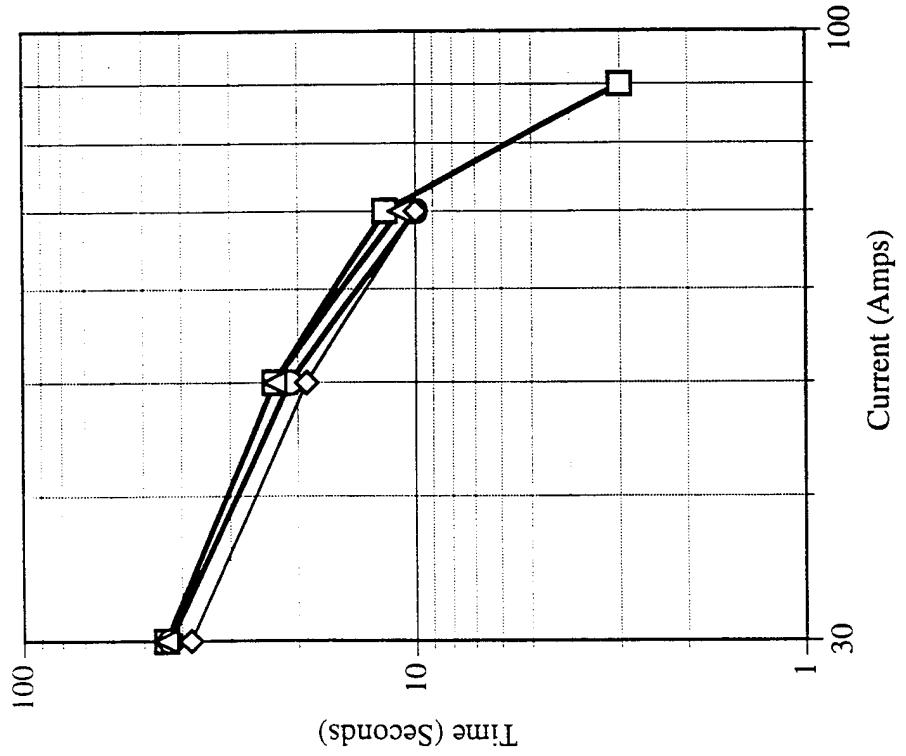


FIGURE 23
Small Scale Characterization
Ragone Relationship, 24 deg C

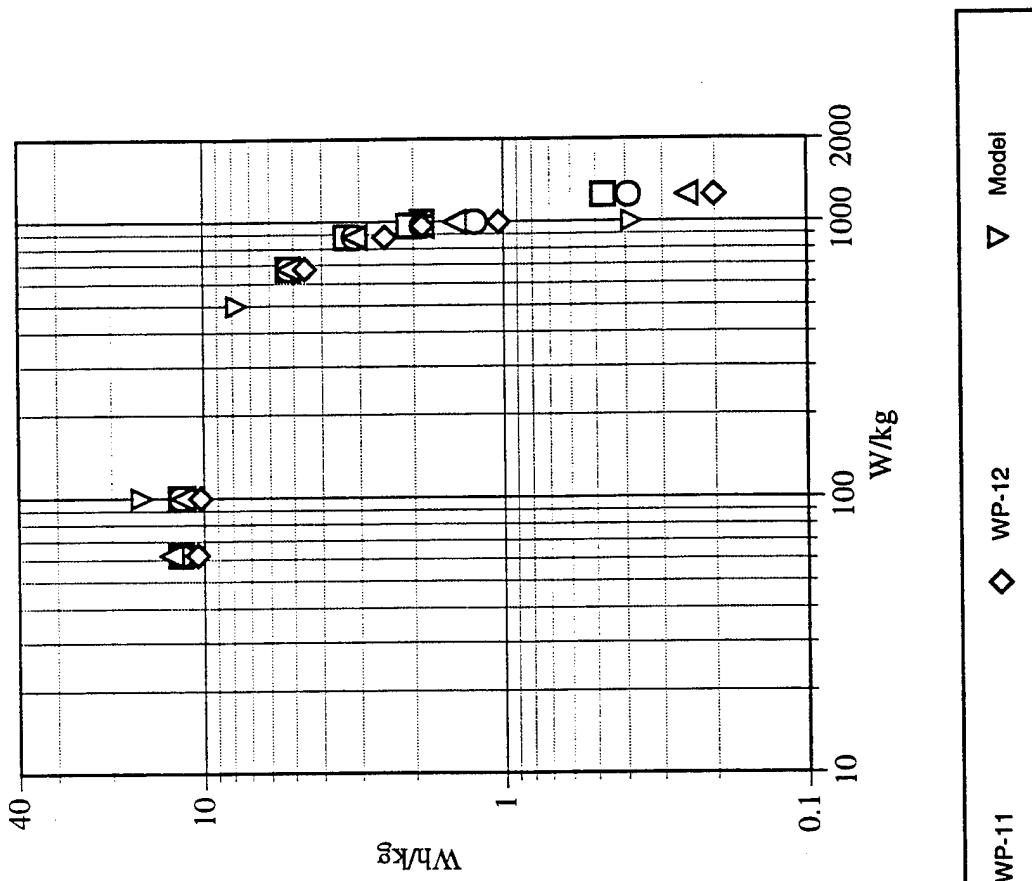
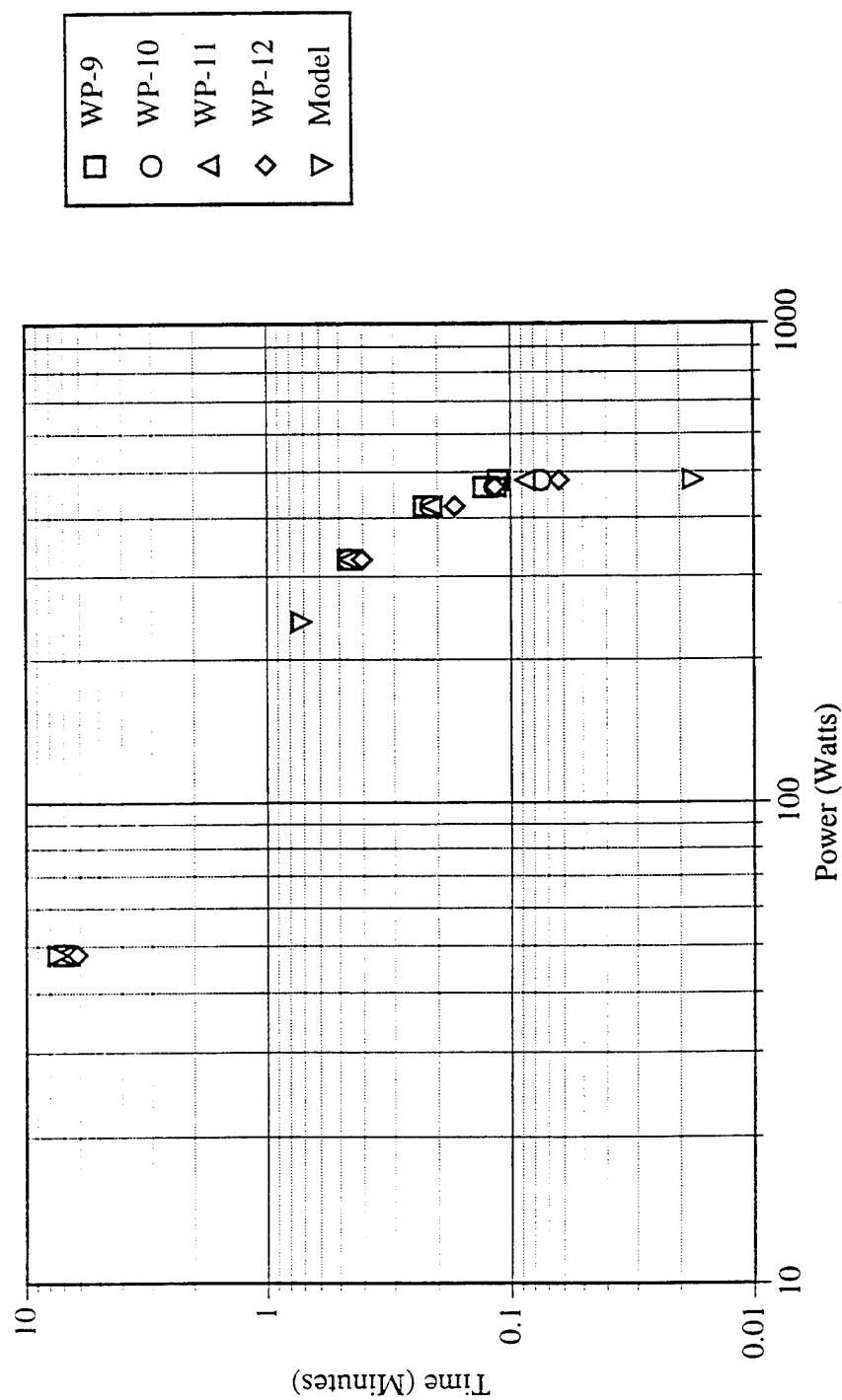


FIGURE 24
Small Scale Characterization
Discharge Time vs Power, 24 deg C



sheet, chilled, slit into ribbon, chilled again, and finally laid onto treated plastic spacer frames. Dummy stacks were leak-free to 4 psig, and successfully completed 4 hours of intermittent vacuum pulsing to simulate the fill and form procedure. Cells in 12- and 24-volt stacks were also leak-free when similarly tested.

Sufficient quantities of the engineering sample material resided in-house, but efforts to find a replacement adhesive were initiated when additional material could no longer be obtained. Chemical analyses and physical testing of the original material was requested of H.B. Fuller and resulted in their furnishing two candidate replacement materials. Stack assembly showed one sample to be tackier and both materials able to withstand the level of vacuum required for filling. No further work with these substitutes was carried out since sufficient adhesive existed to complete the contract.

4.5.2 Subtask 5.2 Formation

Two 24-volt batteries were stacked for formation studies. Each was comprised of 13 electrodes, 12 spacer frames, and two copper termination plates bolted between polycarbonate endwalls and outfitted with polycarbonate filling manifolds across each set of top slots.

Air pressurization of the first stack prior to filling showed one of the two fill ports to be leaking. The manifold was removed and the slots closed off after repeated attempts to seal the manifold were unsuccessful. Seventy-five minutes were required to input 300 cc of chilled electrolyte through the remaining manifold. This represented roughly 72% of the available void volume in the stack. Complete (100%) saturation had been targeted, however, small leaks developed around the base of the manifold, decreasing the fill efficiency. Current was applied for 120 minutes when evidence of shorting was apparent. Disassembly showed the majority of cells to have dendritic shorting through the center area of the separator. Failure was attributed to the long fill time (10-15 minutes was targeted to minimize the dissolution and diffusion of lead into the separator) and out-of-spec plate thicknesses. On average, plates were 0.007" over the 0.025" target, resulting in a compressed separator allowance of 0.016". Roughly 0.020" was considered the minimum separator thickness. Paste weights were reduced for the subsequent build.

The second 24-volt battery was assembled into a bolted polycarbonate fixture, filled to 84% saturation with chilled electrolyte, and placed on formation. Further filling risked lead dissolution and dendrite formation in the separator due to the excessive time required. Five cells shorted during formation as a result of a common electrolyte path along the lead exposed within the fill channel. Further formation attempts were placed on hold pending receipt of a molded stack which, by design, better guarded against common electrolyte paths in the fill port area.

4.6 WBS 6.0 BMET DEMONSTRATION

4.6.1 Subtask 6.1 Deliverables

Injection molded containment about metallic substrates was aggressively pursued for the majority of the No-Cost Time Extension. Repeated trials ultimately succeeded in correcting recurrent frame and electrode distortion, however, hermetic cell-to-cell seals were not obtained. Stacks were never available for formation or for trials to attach covers via induction welding. As a result, a backup battery design was implemented to complete the contract's deliverable requirements.

The following section describes the injection molded containment work in more detail, along with the proposed venting and intermodule connector concepts. The subtask is then concluded with a description of the batteries delivered to WPAFB.

4.6.1.2 24-Volt Injection Molded Containment

The use of the injection molded containment concept previously tested with composite electrodes required one design modification to facilitate use with metallic substrates. To prevent distortion of the 0.012" thick metal electrode, the outer edge of the spacer frame was reshaped to wrap around the lead sheet and afford protection against the injection pressure. Glass filler was also added to the spacer resin to promote a melt bond with the outer endwalls. Molded spacers showed that shrinkage of the 0.082" thick parts was less than anticipated (0.003 in/in vs. 0.007 in/in). This was due to the ASTM shrinkage rate reporting basis (0.125"x0.5"x6" sample). As a result, spacers were slightly larger than specified, however, down-the-line assembly problems were not encountered.

The endwall material was also reevaluated and three candidates tested for use in maintaining the compressed stack dimension. Single layers of honeycombed aluminum sheet stock failed deflection testing. Bulk molding compound manufactured by Luvdahl provided the needed strength against a 6 psig load but was incompatible with battery acid. Glass-filled polypropylene was ultimately used after measuring a deflection of 0.013" at 5 psig.

Severely warped endwalls were produced during the first mold trial. Mold gate changes reduced the distortion, but a subsequent heat soak was still necessary to produce a flat part. Limited success was had in adding a blowing agent. Topical sinks located around the outer perimeter and the center termination port were greatly reduced but not eliminated. Slight part warpage also remained. Cross sectioning showed the internal pore size (caused by the blowing agent) to be very small. It also showed a 4-hour heat treatment to cure the warpage with no sign

of reactivating the blowing agent, but at the expense of the recessed terminal electrode cavity dimension. Heat treating was abandoned when measurements showed shrinkage along the length and width centerlines to be so great as to make it impossible to insert the terminal electrode in the recessed cavity.

Endwalls and spacers were then assembled with lead sheet to create dummy stacks for mold trials. Early attempts showed the plastic to distort the 10% glass frames inward toward the pasted portion of the stack, leaving insufficient material to fill the outer frame. Gate modifications were implemented in an effort to equalize the injection pressures at various points within the container mold. Center/side gating achieved complete mold fill and eliminated much of the frame distortion, however, cross-sectioning still showed buckled lead and uneven plastic distribution. The mold clamp location was then widened and additional glass added to the spacer resin for strength.

Strengthened plastic battery components were received and set up parts prepared for a trial in mid-July, 1995. Glass loading in the frame was increased to 30% in order to prevent blowing in and lead distortion, and to reduce part compression when clamped within the mold. The molding trial was nearly successful. Complete mold fill was achieved with slight crowning of the frames. A "clamp only" trial showed the crowning to be a result of the mold closing. Still closer examination revealed the stacks loaded into the mold to be ~0.100" too thick as a result of out-of-spec adhesive. The remaining thick stacks were preheated and easily compressed to the correct 1.454" thick dimension. Disassembly showed no electrode distortion. Laboratory measurements of stacks assembled using 0.003" thick adhesive (a 50% reduction) were similarly flat.

The subsequent molding trial with correctly processed adhesive produced four dummy stacks and one DUF battery for analysis. Electrodes in all four dummy stacks were distorted along the inner frame perimeter. Heat sensitive indicators inserted at two points in each stack recorded the temperature history and showed no indication of having reached the temperature at which the inlaid adhesive would begin to flow.

The distortion was subsequently eliminated in late July by thermally fusing the outer edges of the stack to better resist the high molding pressure. Pressure testing to confirm cell-to-cell seals identified leakage that was traced to the area surrounding the fill channels. Close examination showed a lack of melt bond between the prefused frame and injected containment plastic. Given the cost and time associated with the mold change proposed to eliminate the leakage, the concept was abandoned for use with WPAFB deliverables.

Venting considerations were evaluated concurrently to stack molding. Implementation of a totally sealed design was initially considered, but dismissed. Utilizing a fail-safe panel along

the face representing the endwall would have reduced its functionality as a means of maintaining adequate battery compression. User safety in the event of an abusive overcharge was an even greater concern.

A review of available off-the-shelf vents quickly showed that no battery vent supplier had ever addressed the main issue facing bipolar technology: cell width. Vent designs just 0.060" to 0.080" in width did not exist. Staggering the vents was proposed, but eliminated from further consideration when it became apparent that multiple frame molds would be required.

Having limited data showing success in cycling a small bipolar battery utilizing single point venting, the deliverable venting configuration was drawn. In its final form, a 24-volt battery was to be fitted with a vent over each of the fill slot locations. This duplicity provided a backup venting location to any cell that might incur blockage in one of its ports. Oil applied topically aided in achieving and maintaining the hermetic seal required for recombinant, maintenance-free operation.

Two methods were suggested for attaching the vent/cover to the injection molded battery housing: heat sealing and induction welding.

Heat sealing is used throughout the battery industry. Generally, this involves heating the edges to be joined, bringing them into contact, and allowing them to cool under pressure. Concern was raised over being able to hold the 0.080" thick cover while preheating it with a heat lamp. That and the estimated \$30,000 to build a suitable machine to try the concept made heat sealing a last choice technique.

Induction welding was then investigated. This process was reportedly fast and versatile. Heat induced by a high frequency electrodynamic field in a metallic insert placed at the joint brings the surrounding material to the melt temperature. Pressure maintained as the field is turned off maintains the joint as it solidifies. Welding occurs only in the area immediately adjacent to the metallic insert. As a result, weld strength depends on the size and geometry of the metal insert.

The process was also feasible economically. Purchasing a new laboratory unit required \$10,000. Leasing was also possible at \$750 per month.

Initial induction welded samples prepared by Pillar Industries indicated that a hermetic bond could be easily achieved around the periphery of the vent/cover. A semicircular cavity rimming the upper edge of the battery and the two cross bars spanning the center portion of the upper surface was included in the mold design. Later testing proved that a hermetic bond along the cross bars would not be achieved. Mold changes were ordered to reduce the cross bar height to make them serve only as structural supports. Hermetic seals at these points were not necessary given the remainder of the cover weld met specification. Test welds with stacks and covers were never attempted given the difficulties previously described.

Lastly, NCTE work was performed to efficiently connect two 24-volt units in series to form a higher voltage subassembly. Various porous copper samples were obtained and tested under load. Results showed the porous copper to be less resistive than solid copper sheet wrapped around a foam pad (Figure 25). Twenty pieces of 60 pores per inch (ppi) material were ordered and received on time, but never used in deliverables. The batteries delivered utilized a backup containment design that facilitated direct assembly of higher voltage stacks.

4.6.1.3 Gasketed Containment

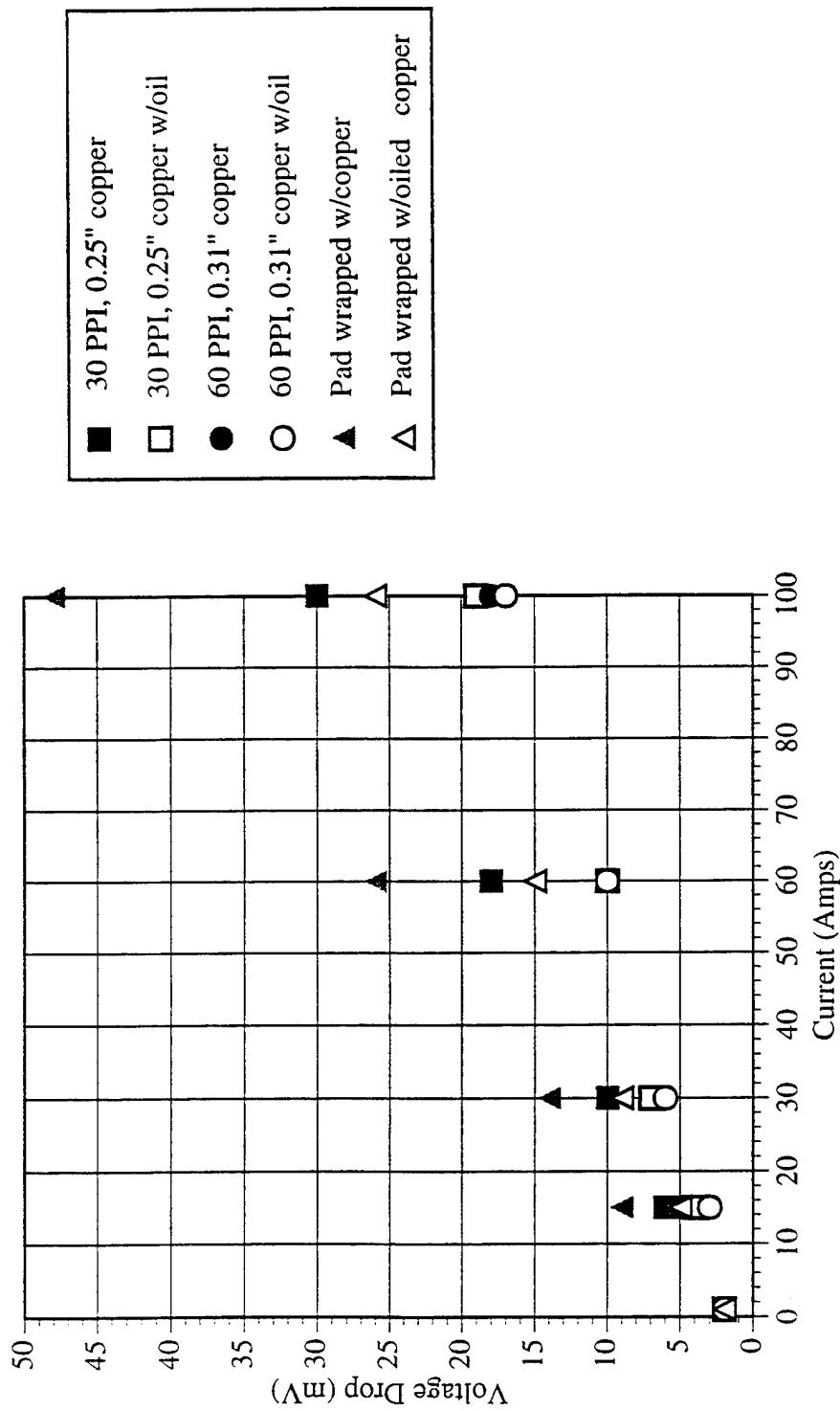
Given the difficulties encountered in achieving hermetic cell-to-cell seals with the injection molded containment concept, WPAFB accepted deliverable batteries assembled using neoprene spacers and machined ABS container components (Appendix B).

Bipolar electrode substrates were die cut from 0.012" thick tin-lead sheet and pasted following the attachment of plastic screen (see Metallic Substrate Development, Subtask 3.3.2). Three paste runs succeeded in pinpointing the wet paste weight needed to achieve the targeted 0.062" electrode thickness. After curing and drying, plates were individually cleaned, weighed, and checked for high spots (thickness). Paste mass and thicknesses of bipolar electrodes used in the deliverable candidates were put at 105.7 ± 2.5 grams and 0.059 ± 0.001 ", respectively.

Terminal electrodes were die cut from laminated sheet stock comprised of 0.008" thick lead and 0.014" thick copper. This design permitted copper terminations to be soldered to the copper face of the electrode with minimal risk of burning a pinhole through the lead face. Each 0.75" long x 0.75"OD stud with a tapped thread was correctly located by first soldering it to an oversized electrode that was then die-cut to achieve the required dead-center location. (This procedure had been critical to injection mold trials since the stack position in the mold was based on the stud location.) Stud welds were shown to withstand an average of 285 in-lb of torque before failing at the solder-to-laminate joint. This compared favorably to the 180 in-lb SLI specification.

Container components were machined from 0.125" and 0.250" (nominal) thick ABS. Solvent bonding was implemented to join the pieces. Endwalls were provided the necessary strength by encapsulating multiple sheets of honeycombed aluminum within a protective ABS cavity. Electrodes were sequentially placed onto neoprene gaskets and absorptive glass mat positioned over the active area to prevent shorting. Separator material was sized to overlap the active area slightly. Starting thickness facilitated the 25% compression deemed critical to supporting high rates of discharge. Fittings were located in channels milled into each gasket to create ports for filling and venting.

FIGURE 25
Voltage Drop Across Intermodule Connector Candidate Materials



Fill and formation were attempted only after confirming each and every cell in a stack to be leak free. Filling was accomplished by evacuating the cells through a column of chilled electrolyte. Returning the system above the electrolyte to atmospheric pressure forced the predetermined volume of acid into each cell quickly and efficiently. Internal stack temperature was monitored constantly and used in controlling the formation current. Current was applied as soon as the fill was completed to minimize the risk of dendritic shorting due to lead dissolution.

Fittings were removed and the cover/vent assembly solvent bonded into place after limited qualification cycling was performed to fully develop the capacity. Details regarding the assembly, formation, and qualification testing of each deliverable are included in Appendix B.

To assist WPAFB in preparing for receipt of these units, three bound copies of safety instructions and operating recommendations were mailed February 29, 1996. One 24-volt and two 12-volt nominal batteries were hand delivered to Wright Laboratory on March 6, 1996 with an additional two copies of the instructions and recommendations. Identification and safety labels were attached to each battery to warn of the potential for explosion, acid burns and electrical shock.

APPENDIX A

RESISTIVITY TESTING

RESISTIVITY TESTING

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
47A	4/2/92	LAMINATED 85% GC23N W/O CA 15% MICROTHERENE 4.5 M.I.	0.365 0.450 0.270	0.023 0.024 0.023	6.248 7.382 4.622	0.960 0.640 1.060	0.042 0.042 0.042	8.999 5.999 9.936	36.6158656 8.311
48A	4/2/92	C-PLASTIC LAMINATED 85% GC23N WITH CA 15% MICROTHERENE 4.5 M.I.	0.630 0.570 0.635	0.025 0.024 0.024	9.921 9.350 10.417	1.000 1.500 0.740	0.040 0.040 0.040	9.843 14.764 7.283	14.764 7.283 7.415184
52	4/9/92	C-PLASTIC LAMINATED 84% GC23N & 16%PTFE TO C-PLASTIC & Pb FOIL SINGLE APPLICATION OF RESIN	1.180	0.053	8.765	19.500 17.500 38.000	0.062 0.062 0.063	123.825 111.125 237.470	1696.53351 157.474
53	4/9/92	C-PLASTIC LAMINATED 84% GC23N & 16%PTFE TO C-PLASTIC & Pb FOIL DOUBLE APPLICATION OF RESIN	3.630	0.061	23.428	169.000 161.000 188.000	0.054 0.054 0.054	1232.138 1173.812 1370.662	5273.26123
54B	4/14/92	LAMINATED 85% GC23N-1 15% MICROTHERENE 4.5 M.I. W/O Pb FOIL	0.195	0.040	1.919	0.440 0.483 0.470	0.040 0.040 0.040	4.331 4.754 4.626	138.119658 4.570
55B	4/14/92	LAMINATED 85% GC23N-2 15% MICROTHERENE 4.5 M.I. W/O Pb FOIL	0.250	0.030	3.281	1.730 2.350 3.600	0.030 0.030 0.030	22.703 30.840 47.244	33.596 924
71A	4/24/92	LAMINATED THICK/THICK GC23N-1 /C-PLASTIC	0.295 0.300 0.280	0.206 0.208 0.208	0.564 0.568 0.530	0.390 0.430 0.400	0.209 0.211 0.208	0.735 0.802 0.757	38.0646797
72A	4/24/92	LAMINATED THIN/THIN GC23N-2 /C-PLASTIC	0.275 0.265 0.275	0.208 0.208 0.207	0.521 0.502 0.523	0.495 0.410 0.500	0.210 0.210 0.209	0.928 0.769 0.942	70.763192
73A	4/24/92	LAMINATED THICK/THIN GC23N-3 /C-PLASTIC	0.420 0.250 0.220	0.031 0.032 0.033	5.334 3.076 2.625	3.800 8.800 6.400	0.031 0.031 0.032	48.260 111.760 78.740	2063.77933 79.587

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
74A	4/24/92	LAMINATED THIN/THIN GC23N-4 /C-PLASTIC	0.380	0.032	4.675	4.300	0.032	52.904	
75A	4/24/92	LAMINATED THICK/THIN GC23N-5 /C-PLASTIC	0.440	0.033	5.249	5.100	0.033	60.845	
			0.430	0.031	5.461	5.500	0.032	67.667	
					5.129			60.472	1079.12921
76A	4/24/92	LAMINATED THIN/THICK GC23N-6 /C-PLASTIC	0.350	0.121	1.139	0.570	0.122	1.839	
			0.580	0.122	1.872	0.520	0.123	1.664	
			0.280	0.123	0.896	0.320	0.123	1.024	
					1.302			1.509	15.9055035
77A	5/12/92	LAMINATED GC23N-A-3/92 Pb-FOIL C-PLASTIC	0.285	0.124	0.905	2.350	0.125	7.402	
			0.275	0.126	0.859	2.580	0.124	8.192	
			0.270	0.126	0.844	2.300	0.123	7.362	
					0.869			7.652	780.246688
78A	6/5/92	LAMINATED GC23N-MICROTHENE & C-PLASTIC	0.36	0.026	5.451				
		LAMINATED GC23N-B-3/92 Pb-FOIL C-PLASTIC	0.22	0.026	3.331				
		LAMINATED GC23N-B-3/92 Pb-FOIL C-PLASTIC	0.66	0.027	9.624				
79A	5/20/92	LAMINATE GC23N-1-85% MICROTHENE/CA	0.52	0.046	4.451	3.7	0.046	31.667	611.538462
		GC23N-2-85% MICROTHENE	0.335	0.044	2.997	2.2	0.044	19.685	556.716418
		GC23N-3-80.3% K _Y	0.49	0.039	4.946	2.1	0.041	201.652	3976.65505
		GC23N-4-80.3%	0.435	0.037	4.629	13.5	0.038	139.867	2921.77858

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
80A	5/27/92 PG.139/141	KY/CA LAMINATE GC23N-1-85% MICROTHENE/CA GC23N-2-85% MICROTHENE GC23N-3-80.3% KY GC23N-4-80.3% KY/CA	0.36 0.495 0.223 0.305	0.04 0.039 0.044 0.042	3.543 4.997 1.995 2.859	0.445 0.525 0.253 0.35	0.04 0.038 0.044 0.041	4.380 5.439 2.264 3.361	23.611111 8.85167464 13.4529148 17.5529788
81A	6/9/92	LAMINATED GC23N/MICROTHENE & C-PLASTIC 5/92-1R 5/92-2R 5/92-3R 5/92-4R							
82A	6/10/92	LAMINATED DOPED OXIDE/SCW AND C-PLASTIC	0.38	0.098	1.527	23.3	0.098	93.604	6031.57895
84A	6/26/92	LAMINATED DOPED OXIDE-5/92 KY 7201 & 711 C-PLASTIC CA	1.7 0.54 0.45 0.785 0.68 0.32	0.031 0.031 0.059 0.032 0.033 0.062	21.590 6.858 9.003 9.658 8.113 0.062	26.3 22.0 1.75 6.4 2.032	0.031 0.033 0.031 0.032 0.032	334.011 2624.672 22.225 78.740	1447.05882 38171.6049 130.121225 870.588235
85A	6/30/92	LAMINATES DOPED OXIDE, CA C-PLASTIC, Pb FOIL 711 KYANR & Pb DUST 70%-W/CA-FOIL 70%-W/O CA-FOIL 70%-W/O CA-DUST 75%-W/CA-FOIL 75%-W/CA-DUST 75%-W/O CA-FOIL	3.7 0.6 2.15 0.32 0.097 0.43 1.25	0.022 0.022 0.022 0.025 0.024 0.024 0.025	66.213 10.737 38.475 5.039 1.591 1.2 71.5 1.2 7.3		0.025 0.025 0.025 0.026	1125.984 188.976 11776.2887 1105.391 15570.8408	22243.75 11776.2887 15570.8408

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
88A	7/13/92	5/92-DOPED OXIDE 75%-W/O CA-DUST	0.3	0.026	4.543	7.0	0.028	984.252	21566.6667
		KY-711 C-PLASTIC	0.11	0.013	3.331	0.115	0.012	3.773	13.2575758
		013-C-PLASTIC	0.65	0.033	7.755				
		020-DOPED OXIDE	1.15	0.042	10.780	3.85	0.042	36.089	234.782609
		030-DOPED OXIDE	1.55	0.05	12.205	1.13	0.048	9.268	-24.0591398
		040-DOPED OXIDE	1.65	0.054	12.030	1.68	0.054	12.248	1.81818182
		050-DOPED OXIDE							
92A	7/22/92	LEAD DUST & POLYSULFONE PREMIXED W/1.1 DRIED & PRESSED AT 599F 30 TONS 55% BY WT.	0.043	0.028	0.605				
	PG. 167								
94A	7/28/92	LAMINATE DOPED OXIDE(5/92) KY-711 & KET WITH KY-711							
		14%-KET/KYN-.050	0.305	0.068	1.766	0.48	0.068	2.779	57.3770492
		14%-KET/KYN-.040	0.38	0.061	2.453	0.4	0.061	2.582	5.26315789
		14%-KET/KYN-.030	0.5	0.051	3.860	0.74	0.051	5.713	48
		Pb/POLYSULFONE	0.066	0.025	1.039	0.57	0.025	8.976	763.636364
95A	7/30/92	LAMINATE DOPED OXIDE(5/92) KY-7201 & KET WITH KY-7201							
		14%-KET/KYN-.050	0.305	0.066	1.819	0.345	0.066	2.058	13.1147541
		14%-KET/KYN-.040	0.295	0.061	1.904	0.4	0.061	2.582	35.5932203
		14%-KET/KYN-.030	0.27	0.052	2.044	1.15	0.052	8.707	325.925926
		14%-KET/KYN-.020	0.255	0.043	2.335	4.2	0.043	36.454	1547.05882
96A	8/10/92	LAMINATES DOPED OXIDE w/MICROTHENE KET & MICROTHENE							
		80%-DOPED OXIDE-96A-1	0.62	0.071	3.438	0.64	0.071	3.549	3.22580645
		80%-DOPED OXIDE-96A-2	0.46	0.063	2.875	0.44	0.063	2.750	-4.34782609

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH)	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
97A	8/18/92	LAMINATES DOPED OXIDE/KY(8/92)	0.21 0.23 0.218	0.052 0.063 0.067	1.590 1.437 1.281	1.65 0.43 0.29	0.052 0.063 0.067	12.492 2.687 1.704	685.714286 86.9565217 33.0275229
99A	8/21/92	LAMINATES DOPED OXIDE/KY KET/KY	0.25 0.22 0.225 0.185	0.074 0.076 0.06 0.06	1.330 1.140 1.476 1.214	0.31 0.32 0.51 0.33	0.074 0.076 0.06 0.06	1.649 1.658 3.346 2.165	24 45.454555 126.66667 78.3783784
102A	9/16/92	LAMINATES DOPED OXIDE/MICROTHENE KET/MICROTHENE	1.55 1.15 1.75 1.13	0.061 0.073 0.049 0.046	10.004 6.202 14.061 9.671	2.4 1.63 3.4 1.83	0.062 0.077 0.049 0.048	15.240 8.334 27.318 15.010	52.3413111 34.3760587 94.2857143 55.199115
103A	9/23/92	LAMINATES WASHED DOPED OXIDE PRECOMPOUNDED C-PLASTIC	0.58 0.595 0.375 0.355	0.08 0.063 0.05 0.04	2.854 3.718 2.953 3.494	1.5 6 2.8 12.5	0.08 0.063 0.05 0.04	7.382 37.495 22.047 123.031	158.62069 908.403561 646.666687 342.112676
104A	9/29/92	LAMINATES WASHED DOPED OXIDE PRECOMPOUNDED C-PLASTIC	0.33 0.44 0.31 0.355 0.72 0.7 0.455 0.54	0.047 0.058 0.064 0.073 0.048 0.062 0.066 0.066	2.764 2.987 1.907 1.915 5.906 4.445 2.714 3.221	8.5 3.2 5.2 2.9 10.3 5.5 5.5 4.3	0.047 0.058 0.064 0.073 0.048 0.062 0.066 0.066	71.201 21.721 31.988 15.640 84.482 34.925 32.808 25.650	2475.75758 627.27277 1577.41935 716.901408 1330.55556 685.714286 1108.79121 696.296296
105A	10/9/92	KY (7/92) & MICROTHENE (5/92)							

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
80%-LOADING									
		DOPED OXIDE (5/92)							
		10%KY/90%MIC.-105A-1	0.17	0.056	1.195	0.45	0.056	3.164	164.705882
		20%KY/80%MIC.-105A-2	0.185	0.053	1.374	0.78	0.053	5.794	321.621622
		30%KY/70%MIC.-105A-3	0.173	0.053	1.285	1.85	0.053	13.742	969.364162
		40%KY/60%MIC.-105A-4	0.165	0.05	1.299	2.8	0.05	22.047	1596.9697
109A	KY (7/92) & MICROTHENE (5/92)	80%-LOADING							
		DOPED OXIDE (5/92)							
		109A-1	0.29	0.041	2.785				
		109A-2	0.36	0.04	3.543	0.87	0.04	8.563	141.666667
		109A-3	0.33	0.041	3.169	4.4	0.042	412.4448	12915.873
		109A-4	0.44	0.039	4.442	0.85	0.041	8.162	83.7583149
110A	10-26-92	80% DOPED OXIDE(5/92)							
		MICRO.(5/92) & KY(7/92)							
		110A-1	0.225	0.038	2.331	4.9	0.038	50.767	2077.777778
		110A-2	0.35	0.039	3.533	4.8	0.039	48.455	1271.42857
		110A-3	0.22	0.042	2.062	1.75	0.042	16.404	695.454545
		110A-4	0.33	0.041	3.169	0.57	0.041	5.473	72.7272727
111A	10/29/92	LAMINATES							
		PRECOMPOUNDED MICRO/DOPED OXIDE							
		85%-LOADING							
		111A-1	1	0.042	9.374	1.95	0.042	18.279	95
		350F/3 TONS							
		80%-LOADING							
		111A-2	2.1	0.043	19.227	3	0.043	27.467	42.8571429
		KY/DOPED OXIDE							
		75%-LOADING							
		111A-3	0.8	0.053	5.943	1.2	0.053	8.914	50
		MICRO/DOPED OXIDE							
		80%-LOADING							
		350F/3 TONS							
		111A-4	1.8	0.036	19.685	2	0.036	21.872	11.111111
112A	11/5/92	LAMINATES							
		75% LOADING							
		DOPED OXIDE(7/92)							
		KY(7/92)							
		14%KET(9/92)							
		400F/3 TONS	112A-1	0.15	0.089	0.664	0.089	0.000	-100
		400F/3 TONS	112A-2	0.165	0.088	0.738	0.088	0.000	-100

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
LAMINATES 80% LOADING									
DOPED OXIDE(7/92) MICROTHENE(5/92) PRECOMPOUNDED C-PLASTIC									
325F/3 TONS 325F/3 TONS									
113A	11/10/92	LAMINATES	0.46	0.069	2.625	0.069	0.000	0.000	-100
80% DOPED OXIDE(7/92) MICROTHENE(5/92) CA PRECOMPOUNDED C-PLASTIC									
325F/3 TONS 325F/3 TONS 325F/3 TONS 325F/3 TONS									
114A	11/10/92	LAMINATES	0.37	0.066	2.207	4.6	0.066	27.440	1143.24324
80% DOPED OXIDE(7/92) MICROTHENE(5/92) CA PRECOMPOUNDED C-PLASTIC									
325F/3 TONS 325F/3 TONS 325F/3 TONS 325F/3 TONS									
115A	11/24/92	LAMINATES	0.36	0.074	1.915	0.85	0.074	4.522	136.11111
80% LOADING DOPED OXIDE 20% MICROTHENE									
WASHING TECH. PRECOMPOUNDED C-PLASTIC .2%/.07GMS CA									
325F/3 TONS 115A-1 COARSE-X									
325F/3 TONS			0.38	0.073	2.049	0.52	0.072	2.843	38.7426901

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
	325F/3 TONS	115A-2 COARSE-X	0.35	0.072	1.914	0.52	0.072	2.843	48.5714286
	325F/3 TONS	115A-3 MEDIUM-X	0.46	0.062	2.921	0.96	0.062	6.096	108.695652
	325F/3 TONS	115A-4 MEDIUM-X	0.58	0.067	3.408	1.18	0.067	6.934	103.448276
116A 11/25/92 80% LOADING DOPED OXIDE									
20% MICROTHENE									
PRECOMPOUNDED C-PLASTIC 2% / 07GMS CA									
325F/3 TONS									
	325F/3 TONS	116A-1 COARSE	0.37	0.077	1.892				
	325F/3 TONS	116A-2 COARSE	0.3	0.071	1.664				
	325F/3 TONS	116A-3 MEDIUM	0.88	0.067	5.171				
	325F/3 TONS	116A-4 MEDIUM	0.61	0.066	3.639				
117A 12/03/92 80% LOADING DOPED OXIDE									
20% MICROTHENE									
PRECOMPOUNDED C-PLASTIC .15% TO .45% CA									
325F/3 TONS									
	117-1A (.15%)	0.38	0.071	2.107	0.98	0.071	5.434	157.894737	
	117-2A (.20%)	0.51	0.071	2.828	1.15	0.071	6.377	125.490196	
	117-3A (.25%)	0.42	0.068	2.432	0.7	0.066	4.176	71.7171717	
	117-4A (.30%)	0.56	0.068	3.242	0.98	0.068	5.674	75	
	117-5A (.35%)	0.42	0.071	2.329					
	117-6A (.40%)	0.46	0.065	2.786					
	117-7A (.45%)	0.64	0.064	3.937					
118A 12/07/92 80% LOADING DOPED OXIDE									
20% MICROTHENE									
PRECOMPOUNDED C-PLASTIC .15% TO .45% CA									
325F/3 TONS									
	118-1A (.15%)	0.4	0.068	2.316					

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
118		118-2A (.20%) 118-3A (.25%) 118-4A (.30%) 118-5A (.35%) 118-6A (.40%) 118-7A (.45%)	0.4 0.38 0.33 0.3 0.4 0.36	0.071 0.068 0.068 0.068 0.069 0.068	2.218 2.200 1.911 1.737 2.282 2.084	0.069 0.067 0.4 0.46 0.45 0.45	2.282 2.350 2.625 0.689 2.605 0.068	19.4554238 35.3233831 15 2.5	
119A	12/04/92	THIN LAMINATES 20% MICROTHENE PRECOMPOUNDED C-PLASTIC .25% CA 325F/3 TONS	0.5 0.34 0.225 0.42	0.038 0.038 0.033 0.03	5.180 3.523 2.684 5.512				
119A	119A	80% LOADING DOPED OXIDE							
120A	12/16/92	HAND COMPOUNDED CARBON PLASTIC .25% CA 120-1A 350F 120-2A 350F 120-3A 375F 120-4A 375F	0.43 0.51 0.38 0.44	0.058 0.061 0.059 0.062	2.919 3.292 2.536 2.794	0.52 0.56	0.059 0.062	3.470 3.556	36.8421053 27.2727273
		PRECOMPOUNDED CARBON PLASTIC 120-5A 120-6A	1.3 2.05	0.056 0.058	9.139 13.915	1.95 2.95	0.056 0.058	13.709 20.024	50 43.902439
121	12/17/92	LAMINATES 80% LOADING DOPED OXIDE .25% CA HAND COMPOUNDED CARBON PLASTIC							
		121-1A 121-2A	0.43 0.43	0.075 0.07	2.257 2.418				
122A	12/17/92	LAMINATES 2.60G KETBLACK 10.37G MICRO (5/92) 325F/15 TONS 0.060" SHIM							

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
		122-1A	0.46	0.051	3.551	1.4	0.051	10.807	204.347826
		122-2A	0.43	0.05	3.386	5.4	0.051	41.686	1131.19015
		122-3A	0.41	0.05	3.228	1.45	0.051	11.193	246.724055
		122-4A	0.43	0.052	3.256	0.82	0.052	6.208	90.6976744
123A	01/04/93	LAMINATES 80% LOADING DOPED OXIDE							
		20% MICROTHERENE HANDCOMPOUNDED C-PLASTIC .30% TO 1.00% CA							
		325F/3 TONS							
		123-1A (.30%)	0.49	0.08	2.411	0.58	0.08	2.854	18.3673469
		123-2A (.35%)	0.36	0.081	1.750	0.37	0.081	1.798	2.77777778
		123-3A (.40%)	0.58	0.08	2.854	0.74	0.081	3.597	26.0110685
		123-4A (.45%)	0.43	0.081	2.090	0.51	0.08	2.510	20.0872093
		123-5A (.50%)	0.44	0.078	2.221				
		123-6A (.55%)	0.65	0.078	3.281				
		123-7A (.60%)	0.62	0.076	3.212				
		123-8A (.65%)	0.6	0.076	3.108				
		123-9A (.70%)	0.66	0.078	3.331				
		123-10A (.75%)	0.6	0.076	3.108				
		123-11A (.80%)	0.9	0.08	4.429				
		123-12A (.85%)	0.68	0.08	3.346				
		123-13A (.90%)	0.6	0.072	3.281				
		123-14A (.95%)	0.52	0.075	2.730				
		123-15A (1.00%)	0.54	0.075	2.835				
124	124A 07-JAN-93	LAMINATES 80% LOADING DOPED OXIDE							
		20% MICROTHERENE HANDCOMPOUNDED C-PLASTIC 1.5% TO 3.0% CA							
		325F/3 TONS							
		124-1A (1.5%)	0.62	0.075	3.255				
		124-2A (2.0%)	0.98	0.077	5.011				
		124-3A (2.5%)	0.83	0.076	4.300				

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
									124-4A (3.0%)
125A	125A 01/12/93	LAMINATES TEMP 230F TO 400F 85% DOPED OXIDE PELLETS HANDCOMPOUNDED	124-4A (3.0%)	0.66	0.077	3.375			
		C-PLASTIC							
		125-1A (300F)	0.41	0.057	2.832				
		125-2A (300F)	0.73	0.057	5.042				
		125-3A (350F)	0.42	0.05	3.307				
		125-4A (350F)	0.59	0.051	4.555				
		125-5A (375F)	0.45	0.051	3.474				
		125-6A (375F)	0.44	0.052	3.331				
		125-7A (400F)	0.39	0.051	3.011				
		125-8A (400F)	0.39	0.051	3.011				
		125-11A (275F)	0.58	0.057	4.006				
		125-12A (275F)	0.38	0.057	2.625				
126	126A 01/14/93	LAMINATES 80% TO 90% LOADING DOPED OXIDE(792) .35% CA							
		SAMPLES 1&2 .30% CA							
		SAMPLES 3-7 HANDCOMPOUNDED							
		C-PLASTIC							
		MICROTHENE (5/92) 325F/3 TONS							
		126-1A (80%)	1.45	0.061	9.358				
		126-2A (80%)	2.85	0.061	18.394				
		126-3A (85%)	0.32	0.08	1.575	0.4	0.08	1.969	25
		126-4A (85%)	0.27	0.076	1.399	0.33	0.076	1.709	22.2222222
		275F/3 TONS							
		126-6A (82.5%)	1.4	0.073	7.550	1.45	0.073	7.820	3.57142857
		126-7A (82.5%)	0.52	0.062	3.302	0.53	0.062	3.366	1.92307692
129A	01/15/93	LAMINATES 85% DOPED OXIDE PELLETS 14% TO 22% KET (9/92)							
		325F/3 TONS							
		129-1A (15%)	0.54	0.05	4.252				
		129-2A (15%)	0.64	0.048	5.249				
		129-3A (16%)	0.55	0.049	4.419				
		129-4A (16%)	0.56	0.049	4.499				

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
130A	01/19/93	LAMINATE 325F/3 TONS	129-5A (16%) 129-6A (18%) 129-7A (22%) 129-8A (22%)	0.75 0.66 0.63 0.38	0.05 0.049 0.051 0.05	5.906 5.303 4.863 2.992			
131A	01/27/93	LAMINATE 325F/3 TONS	131-1A(3 TONS) 131-3A(15 TONS) 131-4A(15 TONS) 131-5A(3 TONS)	0.58 0.74 0.51 0.78	0.061 0.055 0.052 0.076	3.743 5.297 3.861 4.041			
132A	01/28/93	LAMINATE 325F/3 TONS	132-1A(3 TONS) 132-2A(3 TONS) 132-3A(15 TONS) 132-4A(15 TONS)	1.3 1.5 0.96 0.79	0.057 0.048 0.05 0.051	8.979 12.303 7.559 6.099			
133A	01/28/93	LAMINATE 325F/3 TONS	133-1A(3 TONS) 133-2A(3 TONS) 133-3A(15 TONS) 133-4A(15 TONS)	0.36 0.32 0.44 0.5	0.069 0.065 0.051 0.052	2.054 1.938 3.397 3.786			
134A	01/28/93	LAMINATE 325F/3 TONS	134-1A(3 TONS) 134-2A(3 TONS) 134-3A(15 TONS) 134-4A(15 TONS)	0.76 0.63 0.85 0.76	0.058 0.057 0.049 0.051	5.159 4.351 6.830 5.867	0.83 0.69 0.72 0.68	0.058 0.058 0.049 0.05	5.634 4.684 5.785 5.354
135A	01/28/93	LAMINATE 325F/3 TONS	135-1A(.010") 135-2A(.010") 135-3A(.006") 135-4A(.006")	0.65 0.6 0.56 0.62	0.029 0.034 0.029 0.029	8.824 6.948 7.602 8.417	0.61 0.57 0.51 0.72	0.03 0.034 0.029 0.029	8.005 6.600 6.924 9.775
136A	02/01/93	LAMINATE 325F/3 TONS	136-1A(22%)	0.39	0.034	4.516			

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		136-2A(22%)	0.57	0.033	6.800				
		136-3A(24%)	0.34	0.035	3.825				
		136-4A(24%)	0.39	0.034	4.516				
137A	02/03/93	LAMINATE 325F/3 TONS							
		137-1A	0.24	0.041	2.305				
		137-2A	0.235	0.043	2.152				
		137-3A	0.305	0.042	2.859				
		137-4A	0.215	0.043	1.969				
MRP	02/03/93	LAMINATE 325F/3 TONS							
		MRP-1	0.48	0.056	3.375	0.59	0.056	4.148	22.9166667
		MRP-2	0.41	0.061	2.646	0.48	0.06	3.150	19.0243902
		MRP-3	0.49	0.054	3.572	0.89	0.054	6.489	81.6326531
		MRP-4	0.43	0.058	2.919	0.47	0.058	3.190	9.30232558
138A	02/04/93	LAMINATE 325F/3 TONS							
		138-1A	0.34	0.03	4.462				
		138-2A	0.6	0.032	7.382				
		138-3A	0.43	0.036	4.703				
		138-4A	0.35	0.035	3.937				
139A	02/05/93	LAMINATE 325F/3 TONS							
		139-1A(18%)	0.39	0.022	6.979				
		139-2A(18%)	0.36	0.025	5.669				
		139-3A(22%)	0.33	0.026	4.997				
		139-4A(22%)	0.4	0.027	4.997				
BR AND R3	02/05/93	LAMINATE 325/3 TONS							
		BR-1	0.76	0.039	7.672				
		BR-2	0.85	0.039	8.581				
		R3-1	0.47	0.042	4.406				
		R3-2	0.51	0.049	4.098				
EXTRUDED 3/24/93		168-1A			1.2				
		100/115/120/125							
		168-2A							
		100/110/120/125							
		168-3A							
		IR TOO HIGH, LAMINATION STOPPED							
169A	03/25/93	LAMINATE 325F/3 TONS							
		169-1A							
		0.89	0.041	8.5461878	>100				

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
170A	03/26/93	LAMINATE 325F/3 TONS							
		170-1A	0.68	0.041	6.5296716	0.66	0.041	6.337622431	-2.94117647
		170-2A	0.9	0.041	8.6422124	0.86	0.041	8.258114077	-4.44444444
171A	03/30/93	LAMINATE 325F/3 TONS							
		171-1A	0.35	0.035	3.9370079	0.74	0.035	8.323959505	111.428571
		171-2A	0.68	0.035	7.6490439	1.05	0.036	11.48293963	50.122549
		171-3A	0.52	0.039	5.2493438				
		171-4A	0.55	0.038	5.6983009				
173A	04/2/93	LAMINATE 325F/3 TONS							
		173-1A	0.34	0.039	3.4322633	0.36	0.04	3.543307087	3.23529412
		173-2A	0.41	0.039	4.1389057	0.48	0.041	4.60917995	11.3622844
175A	04/05/93	LAMINATE 325F/3 TONS							
		175-1A(160)	0.55	0.041	5.281352	0.79	0.041	7.585942001	43.6363636
		175-2A(160)	0.39	0.042	3.655793	0.53	0.042	4.968128984	35.8974359
		175-3A(180)	0.47	0.043	4.3032412	0.68	0.043	6.22596594	44.6808511
		175-4A(180)	0.44	0.042	4.1244844	0.58	0.043	5.310382714	28.7526427
176A	04/06/93	LAMINATE 325F/3 TONS							
		176-1A(160)	0.42	0.04	4.1338883	0.43	0.04	4.232283465	2.38095238
		176-2A(160)	0.49	0.04	4.8228846	0.49	0.041	4.705204532	-2.43902439
		176-3A(180)	0.38	0.039	3.836059	0.39	0.039	3.937007874	2.63157895
		176-4A(180)	0.35	0.038	3.6261915	0.36	0.04	3.543307087	-2.28571429
		176-1A	0.42	0.04	4.1338883	0.67	0.04	6.594488189	59.5238095
		176-3A	0.38	0.039	3.836059	0.59	0.04	5.807086614	51.3815789
		176-4A	0.35	0.038	3.6261915	0.5	0.04	4.921259843	35.7142857

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177A	04/12/93	*176-3A *SAMPLE TESTED FOR 30 DAYS 176-3A READING TAKEN AFTER 1 DAY 176-3A READING TAKEN AFTER 2 DAYS	0.38 0.38 0.38	0.039 0.039 0.039	4.134 4.134 4.134	0.9 0.64 0.833	0.9 0.041 0.04	8.858267717 6.46073087 8.198818898	114.278368 56.282798 98.3265336
177A	04/14/93	LAMINATE 325F/3 TONS	177-1A(160) 177-2A(160)	0.61 0.81	0.041 0.044	5.8574995 7.2476736	0.53 0.74	0.041 0.042	5.089302862 6.936632921
177A	04/14/93	LAMINATE 325F/3 TONS	177-3A(180) 177-4A(180)	1.05 0.84	0.043 0.044	9.6136239 7.5161059	0.77 0.65	0.042 0.043	7.217847769 5.951290972
178A	04/14/93	LAMINATE 325F/3 TONS	178-1A(160) 178-2A(160)	0.54 0.64	0.046 0.047	4.6217049 5.361032	0.58 0.68	0.046 0.045	4.964053406 5.949255343
178A	04/15/93	LAMINATE 325F/3 TONS	178-3A(180) 178-4A(180)	0.53 0.45	0.045 0.041	4.6369204 4.3211062	0.48 0.48	0.045 0.041	4.199475066 4.60917995
179A	04/15/93	LAMINATE 325F/3 TONS	179-1A(160) 179-2A(160)	0.39 0.31	0.045 0.043	3.4120735 2.838308	0.46 0.39	0.045 0.043	4.024496938 3.570774583
179A	04/15/93	LAMINATE 325F/3 TONS	179-3A(180) 179-4A(180)	0.28 0.31	0.043 0.043	2.563333 2.838308	0.34 0.38	0.043 0.043	3.11298297 3.479216261
181A	04/28/93	LAMINATE 325F/3 TONS	181-1A(200) 181-2A(200)	0.47 0.4	0.063 0.059	2.9371329 2.6691579	0.58 0.56	0.062 0.057	3.683007366 3.86793756
181A	04/28/93	LAMINATE 325F/3 TONS	181-3A(180)	0.54	0.064	3.3218504	0.61	0.064	3.75246063

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182A 4V BATTERIES FOR PASTE ADHESION	04/28/93	LAMINATE 325F/3 TONS							
182-1A(200) SANDED		0.68	0.073	3.6673498					
182-2A(200) Pb THEN SANDED		0.7	0.06	4.5931759					
182-3A(180) SANDED		0.68	0.071	3.7706554					
182-4A(180) Pb THEN SANDED		0.6	0.058	4.0727668					
183A 04/29/93	LAMINATE 325F/3 TONS	IR at corner.							
183-1A		0.38	0.043	3.4792163	0.54	0.044	4.831782391	38.8755981	
183-2A		0.38	0.043	3.4792163	0.55	0.048	4.511154856	29.6600877	
183-3A		0.38	0.059	2.5357	0.55	0.059	3.670092086	44.7368421	
183-4A		0.38	0.057	2.6246719	0.58	0.059	3.870278927	47.4576271	
184A 05/04/93	LAMINATE 325F/3 TONS								
184-1A		0.58	0.046	4.9640534	0.78	0.046	6.67579596	34.4827586	
184-2A		0.5	0.046	4.2793564	0.8	0.047	6.7012899998	56.5937447	
184-3A		0.58	0.05	4.5669291	0.76	0.051	5.866913695	28.4651792	
185A 05/05/93	LAMINATE 325F/3 TONS	THICK SUBSTRATE							
185-1A		0.38	0.055	2.7201145	0.58	0.055	4.151753758	52.6315789	
185-2A		0.29	0.048	2.3786089	0.41	0.05	3.228346457	36.7241379	
186A 05/05/93	LAMINATE 325F/3 TONS								
186-1A		1	0.041	9.6024582	1.55	0.042	14.52943382	51.3095238	
186-2A		0.82	0.041	7.8740157	1.3	0.043	11.90258194	51.16227907	
187A	LAMINATE 330F/2 TONS								
187-1A		0.155	0.03	2.0341207	10.5	0.03	137.7952756	6674.19355	

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		187-2A	0.135	0.029	1.832745	20	0.029	271.5177844	14714.8148
		187-3A	0.135	0.029	1.832745	6.2	0.029	84.17051317	4492.59259
		187-4A	0.155	0.03	2.0341207	8.1	0.03	106.2992126	5125.80645
188A		LAMINATE 330F/2 TONS							
		188-1A	0.14	0.028	1.9685039	6	0.03	78.74015748	3900
		188-2A	0.14	0.031	1.7780036	9	0.03	118.1102362	6542.85714
		188-3A	0.135	0.031	1.7145034	8.1	0.031	102.8702057	5900
		188-4A	0.155	0.031	1.9685039	9.6	0.031	121.9202438	6093.54839
189A	06/14/93	LAMINATE 295F/3 TONS							
06/14/93		IR OF SAMPLE WAS TOO HIGH. NEW SAMPLES WILL BE MADE AND TESTED							
		189-3A(SANDED)	0.47	0.051	3.6282229	0.7	0.051	5.403736298	48.9361702
		189-4A(SANDED)	0.54	0.045	4.7244094	0.83	0.045	7.261592301	53.7037037
190A	06/16/93	LAMINATE 295F/3 TONS							
		190-1A	0.74	0.086	3.3876579				
		190-2A	0.78	0.092	3.337898				
		190-3A	0.73	0.086	3.3418788				
		190-4A	0.66	0.086	3.0214246				
191A	06/18/93	LAMINATE 295F/3 TONS							
		191-1A(006)SANDED	0.25	0.044	2.2369363	1.6	0.044	14.31639227	540
		191-2A(006)	0.255	0.045	2.2309711	0.97	0.044	8.679312813	289.037433
		191-2A	0.255	0.045	2.2309711	0.38	0.044	3.400143164	52.4064171
		READING TAKEN AFTER BEING STORED FOR 25 MONTHS							
		191-3A(007)SANDED	0.23	0.043	2.1058414	0.48	0.044	4.294917681	103.952569
		191-4A(007)	0.29	0.044	2.5948461	0.58	0.044	5.189692198	100
192A	06/18/93	LAMINATE 295F/3 TONS							
		192-1A	1.3	0.051	10.03551	3.5	0.049	28.12148481	180.21978
		192-2A	1.9	0.049	15.265949	4.4	0.05	34.64566929	126.947368
193A	06/18/93	LAMINATE 295F/3 TONS							
		193-1A(SANDED)	0.19	0.062	1.2065024	0.82	0.062	5.207010414	331.578947
		193-2A	0.26	0.058	1.7648656	2.2	0.059	14.68036834	731.812256

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194A	06/24/93	LAMINATE 295F/3 TONS							
		194-1A(.008")	0.265	0.031	3.3655067	8.5	0.032	104.5767717	3007.31132
		194-2A(.008")	0.32	0.042	2.999625	>10.0	0.043		
		194-3A(.010")	0.29	0.04	2.8543307	>10.0	0.04		
		194-4A(.010")	0.31	0.034	3.5896248	4.4	0.034	509.4951366	14093.5484
195A	06/28/93	LAMINATE 295F/3 TONS							
		195-1A	0.46	0.046	3.9370079	0.65	0.046	5.5631633	41.3043478
		195-2A	0.58	0.046	4.9640534	0.72	0.046	6.162273194	24.137931
196A	06/28/93	LAMINATE 295F/3 TONS							
		196-1A	1.15	0.044	10.289907	1.15	0.045	10.06124234	-2.22222222
		196-2A	1.05	0.045	9.1863517	1.3	0.045	11.3735783	23.8095238
197A	06/29/93	LAMINATE 295F/3 TONS							
		197-1A(315F)	0.24	0.044	2.1474588	0.7	0.045	6.124234471	185.185185
		197-2A(315F)	0.275	0.045	2.40589493	0.65	0.045	5.686789151	136.363636
		197-3A(335F)	0.225	0.045	1.9688039	0.73	0.045	6.386701682	224.444444
		197-4A(335F)	0.36	0.051	2.7790644	0.81	0.051	6.252894859	125
		197-5A(365F)	0.24	0.044	2.1474588	0.37	0.045	3.237095363	50.7407407
		197-6A(335F)	0.22	0.045	1.9247594	0.275	0.045	2.405949256	25
		197-7A(375F)	0.235	0.045	2.055993	0.29	0.045	2.537182852	23.4042553
		197-8A(375F)	0.215	0.045	1.8810149	0.3	0.045	2.624671916	39.5348837
		197-9A(400F)	0.205	0.045	1.7933258	0.275	0.045	2.405949256	34.1463415
		197-10A(400F)	0.2	0.046	1.7117426	0.275	0.045	2.405949256	40.5555556
198A	06/29/93	LAMINATE 295F/3TONS							
		198-1A(315F)	0.43	0.049	3.4549253	0.86	0.049	6.909850554	100
		198-2A(315F)	0.41	0.05	3.2283365	1.25	0.05	9.842519685	204.878049
		198-3A(335F)	0.295	0.047	2.4711007	0.82	0.047	6.868822248	177.966102
		198-4A(335F)	0.32	0.052	2.4222741	0.54	0.052	4.088431254	68.75
		198-5A(355F)	0.28	0.046	2.3964396	1.75	0.044	15.658554044	553.409091
		198-6A(355F)	0.23	0.046	1.9688039	4	0.044	35.7908067	1718.18182
		198-7A(375F)	0.21	0.047	1.7590886	1.95	0.046	16.6894899	848.757764
		198-8A(375F)	0.36	0.049	2.8924456	0.65	0.048	5.331364829	84.3171296
		198-9A(400F)	0.245	0.048	2.0095144	1.2	0.046	10.27045532	411.091393

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
199A									
	07/21/93	LAMINATE 295F/3 TONS 199-1A(NOT SANDED)	0.66	0.044	5.9055118				
		199-2A(NOT SANDED)	0.62	0.043	5.676616				
200A									
	07/23/93	LAMINATE 295F/3 TONS 199-3A(SANDED)	0.31	0.044	2.773801	0.5	0.045	4.374453193	57.7060932
		199-4A(SANDED)	0.37	0.043	3.3876579	0.54	0.044	4.831782391	42.6289926
		LAMINATE 295F/3 TONS 200-1(SANDED)	0.45	0.043	4.1201245	0.58	0.044	5.189692198	25.959596
		200-2(SANDED)	0.47	0.045	4.111936	0.66	0.044	5.905511811	43.6170213
201A									
	07/23/93	LAMINATE 295F/3 TONS 201-1(325)	0.32	0.046	2.7387881	0.4	0.043	3.662332906	33.7209302
		201-2(350)	0.295	0.043	2.7009905	0.4	0.043	3.662332906	35.593203
		201-3(375)	0.34	0.044	3.0422334	0.45	0.045	3.937007874	29.4117647
		201-4(400)	0.31	0.044	2.773801	0.58	0.044	5.189692198	87.0967742
		201-5(425)	0.31	0.044	2.773801	1.4	0.044	12.52684324	351.612903
202A									
	07/23/93	LAMINATE 295F/3 TONS 202-1(325)	0.41	0.043	3.7538912	0.71	0.043	6.500640908	73.1707317
		202-2(350)	0.31	0.043	2.838308	0.48	0.043	4.394799487	54.9387097
		202-3(375)	0.35	0.044	3.1317108	0.54	0.044	4.831782391	54.2857143
		202-4(400)	0.38	0.043	3.4792163	0.54	0.044	4.831782391	38.8755981
		202-5(425)	0.295	0.044	2.6395848	0.98	0.044	8.768790265	232.20339
203A									
	07/23/93	LAMINATE 295F/3 TONS 203-1(325)	0.34	0.044	3.0422334	3.1	0.042	29.05886764	855.182073
		203-2(350)	0.42	0.044	3.758053	2.2	0.044	19.68503937	423.809524
		203-3(375)	0.36	0.044	3.2211883	5	0.042	46.86914136	1355.02646
		203-4(400)	0.5	0.044	4.4738726	3	0.043	27.4674968	513.953488
204A									
	07/27/93	LAMINATE 300F/3 TONS 204-1A(250)	0.62	0.046	5.3064019	1.6	0.046	13.69394043	158.064516
		204-2A(275)	0.45	0.044	4.0264853	0.83	0.044	7.42662849	84.444444
		204-3A(300)	0.51	0.045	4.4619423	0.68	0.045	5.949256343	33.333333
		204-4A(325)	0.51	0.045	4.4619423	0.98	0.045	8.573928259	92.1568627
		204-5A(350)	0.47	0.044	4.2054402	2.9	0.044	25.94846099	517.021277
		204-6A(375)	0.46	0.044	4.1159628	2.35	0.045	20.56993001	399.516908

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
205A									
SEE BATTERY BUILD									
206A	8/10/93	LAMINATE 300F/3 TONS							
		206-1A(SP006)	0.275	0.053	2.0427871	0.34	0.053	2.525627693	23.6363636
		206-2A(SP006)	0.25	0.052	1.8927922	0.31	0.053	2.30277819	21.6603774
		206-3A(SP007)	0.36	0.052	2.7256208	0.48	0.052	3.634161114	33.3333333
		206-4A(SP007)	0.23	0.053	1.7085129	0.34	0.052	2.574197456	50.6688963
207A	8/10/93	LAMINATE 300F/3 TONS							
		207-1A(SP006)	0.83	0.043	7.5993408	1.65	0.042	15.46681665	103.528399
		207-2A(SP006)	0.48	0.041	4.60918	0.64	0.042	5.999250094	30.1587302
		207-3A(SP007)	0.96	0.043	8.789599	1	0.044	8.947745168	1.79924242
		207-4A(SP007)	0.89	0.042	8.34427072	0.86	0.041	8.258114077	-1.01397643
208A	8/11/93	LAMINATE 300F/3 TONS							
		208-1A	0.4	0.037	4.2562247	0.6	0.039	6.056935191	42.3076923
		208-2A	0.5	0.038	5.1802735	0.7	0.038	7.25238226	40
		208-3A	0.3	0.036	3.2809399	0.54	0.037	5.745903584	75.1351351
		208-4A	0.32	0.038	3.3153151	0.73	0.039	7.369271149	122.275641
209A	8/16/93	LAMINATE 300F/3 TONS							
		209-1A(SANDED)	0.96	0.051	7.4108384	3.4	0.051	26.24671916	254.166667
		209-2A	1.35	0.053	10.028228	4.3	0.053	31.941762	218.518519
210A	8/24/93	LAMINATE 300F/3 TONS							
RIBBON FROM DE WAL		210-1A	0.28	0.033	3.3404915	13.75	0.033	164.0419948	4810.71429
		210-2A	0.23	0.033	2.743952	11	0.034	127.3737842	4541.94373
		210-3A	0.41	0.041	3.937079	2.4	0.043	21.97399744	458.139535
		210-4A	0.56	0.042	5.2493438	24	0.043	219.7399744	4086.04651
211A	9/2/93	LAMINATE 300F/3 TONS							

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
212A	9/8/93	LAMINATE 300F/3 TONS	0.56	0.032	6.8897638	0.69	0.03	9.05511811	31.4285714
		212-1A	0.41	0.031	5.2070104	0.62	0.031	7.874015748	51.2195122
		212-2A	0.32	0.024	5.2493438	0.56	0.024	9.186351706	75
		212-3A	0.33	0.023	5.6487504	0.8	0.024	13.12335958	132.323232
213A	9/16/93	LAMINATE 350F/3 TONS	0.86	0.044	7.6950608	1.25	0.044	11.18468146	45.3488372
		213-1A	0.99	0.044	8.8582677	4.4	0.044	39.37007874	344.444444
		213-2A	0.64	0.043	5.8597326	2.3	0.043	21.05841421	259.375
		213-3A	0.72	0.043	6.5921992	1.9	0.043	17.3960813	163.888889
214A	9/20/93	LAMINATE 350F/3 TONS	2.6	0.035	29.246344				
		214-1A	3.4	0.036	37.182852				
		214-2A	3.8	0.035	42.744657				
		214-3A	2.8	0.036	30.621172				
		214-4A	2.5	0.04	24.606299				
		214-5A	3	0.036	32.808399				
215A	9/22/93	LAMINATE 350F/3 TONS	0.5	0.067	2.9380666				
		215-1A	0.6	0.082	2.8807375				
		215-2A	0.84	0.081	4.082823				
		215-3A	0.82	0.081	3.9856129				
		*215-4A	0.86	0.081	4.1800331				
		*215-5A							
216A	9/27/93	LAMINATE 350F/3 TONS							
		LAMINATE 350F/30 TONS							
		216-1A	0.45	0.022	8.0529707	8.3	0.022	148.5325698	1744.44444
		216-2A	0.3	0.021	5.624297	6.9	0.022	123.4788833	2095.45455
		216-3A	0.34	0.022	6.0844667	9.5	0.022	170.0071582	2694.11765
		216-4A	0.36	0.022	6.4423765	1.6	0.022	286.3278454	4344.44444

*SAMPLES NOT SURFACE TREATED

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217A	9/29/93	LAMINATE 300F/3 TONS	217-1A 0.48	0.052	3.6341611	0.66	0.052	4.996971532	37.5
		217-2A 0.43	0.05	3.3858268	0.5	0.05	3.937007874	16.2790698	
		217-3A 0.47	0.052	3.5584494	0.51	0.052	3.867296184	8.5106383	
		217-4A 0.46	0.05	3.6220472	0.6	0.05	4.724409449	30.4347826	
218A	9/29/93	LAMINATE 300F/3 TONS	218-1A 0.9	0.051	6.947661				
		218-2A 1	0.051	7.7196233					
		218-3A 0.7	0.051	5.4037363					
		218-4A 0.73	0.051	5.635335					
219A	10/4/93	LAMINATE 350F/3 TONS							
		LAMINATE 350F/30 TONS							
		219-1A(30 TONS) 0.46	0.038	4.7658516	0.73	0.038	7.563199337	58.6956622	
		219-2A(30 TONS) 0.4	0.038	4.1442188	0.74	0.038	7.666804807	85	
		219-3A(3 TONS) 0.37	0.039	3.73511	0.8	0.04	7.874015748	110.810811	
		219-4A(3 TONS) 0.43	0.04	4.2322835	0.77	0.04	7.578740157	79.0697674	
220A	10/6/93	LAMINATE 300F/3 TONS							
		220-1A 0.34	0.021	6.3742032	0.88	0.022	15.7480315	147.058824	
		220-2A 0.3	0.019	6.2163282	0.69	0.019	14.29755491	130	
		220-3A 0.28	0.02	5.511811	0.54	0.02	10.62992126	92.8571429	
		220-4A 0.34	0.019	7.045172	0.7	0.019	14.50476585	105.882353	
221A	10/11/93	LAMINATE 300F/3 TONS							
		221-1A 0.83	0.045	7.2615923	1.15	0.045	10.06124234	38.5542169	
		221-2A 0.81	0.044	7.2476736	1.1	0.044	9.842519885	35.8024691	
		221-3A 0.85	0.045	7.4365704	1	0.045	8.748906387	17.6470588	
		221-4A 0.92	0.044	8.2319256	1.3	0.044	11.63206872	41.3043478	
222A	10/15/93	LAMINATE 300F/3 TONS							
		222-1A 0.8	0.026	12.111387	1.15	0.026	17.41368867	43.75	
		222-2A 1.2	0.025	18.897638	1.1	0.025	17.32283466	-8.33333333	
		222-3A 0.78	0.025	12.283465	1.15	0.025	18.11023622	47.4358974	
		222-4A 0.91	0.025	14.330709	0.82	0.025	12.91338583	-9.89010989	
223A	10/18/93	LAMINATE 300F/3 TONS							
		223-1A 0.54	0.044	4.8317824	0.55	0.045	4.811898513	-0.41152263	
		223-2A 0.49	0.044	4.3843351	0.7	0.044	6.263421618	42.8571429	
		223-3A(WI) .470/.620	0.044	5.54	1.15	0.045	10.06124234	81.6108726	
		223-4A(WI) .440/.450	0.045	3.93	1	0.044	8.947745168	127.677994	
224A	10/19/93	FIRST WITHOUT SCW, SECOND WITH LAMINATE 350F/3 TONS	224-1A	1.7	0.08	8.3661417			

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*SAMPLES SURFACE TREATED									
225A	10/20/93	LAMINATE 300F/3 TONS (TTS)225-1A (TTS)225-2A (138S)225-3A (138S)225-4A	0.175 0.195 0.235 0.21	1.65 1.7 1.65 1.75 1.85 1.8 1.8 1.8	0.08 0.08 0.08 0.08 0.08 0.08 0.08 0.08	8.1200787 8.3661417 8.1200787 7.7768057 8.6122047 9.1043307 8.8582877 9.7210071 8.7489064	0.36 0.46 0.46 0.285 0.34 0.285 0.28 0.28	0.045 0.044 0.044 0.045 0.045 0.046 0.046 0.046	3.149606299 4.115962777 4.115962777 2.49343832 23.9716332 14.1304348
226A	10/21/93	LAMINATE 300F/3 TONS (TTS)226-1A (TTS)226-2A (138S)226-3A (138S)226-4A (30 DAYS)226-3A	0.14 0.16 0.23 0.28 0.23	0.028 0.028 0.028 0.027 0.028	1.9685039 2.2497188 3.2339708 4.082823 3.2339708	100 100 0.23 0.34 0.28	0.028 0.028 0.028 0.029 0.028	1.406.074241 1.406.074241 3.233970754 4.615802935 3.937007874	71328.5714 62400 0 13.0541872 21.7391304
227A	10/22/93	LAMINATE 300F/3 TONS 227-1A(TTS) 227-2A 227-3A 227-4A	0.84 0.96 0.94 0.94	0.044 0.043 0.044 0.043	7.5161059 8.788599 8.4108805 8.6064823	0.9 1.15 1.1 1	0.044 0.043 0.044 0.045	8.052970651 10.5220271 9.842519655 8.748906387	7.14285714 19.7916667 17.0212766 1.65484634
228A	10/25/93	LAMINATE 300F/3 TONS 228-1A(TTS) 228-1A(TTS) 228-3A(138S) 228-4A(138S)	0.35 0.3 0.47 0.44	0.046 0.045 0.045 0.046	2.9955495 2.6246719 4.111986 3.8495188	0.73 0.67 0.62 0.54	0.046 0.045 0.045 0.045	6.247860322 5.861767279 5.42432196 4.724409449	108.571429 123.333333 31.9148936 22.727277
229A	10/26/93	LAMINATE 300F/3 TONS 229-1A(TTS) 229-1A(TTS) 229-3A(138S) 229-4A(138S)	0.47 0.57 0.74 0.6	0.045 0.045 0.044 0.044	4.111986 4.9868766 6.6213314 5.3686471	0.68 0.74 0.92 0.75	0.044 0.045 0.044 0.044	6.084466714 6.474190726 8.231925555 6.710808876	47.9690522 29.8245614 24.3243243 25

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
230A	10/29/93	LAMINATE 300F/3 TONS							
		230-1A(TTS)	0.29	0.044	2.55948461	0.68	0.044	6.084466714	134.482759
		230-1A(TTS)	0.31	0.043	2.838308	0.59	0.044	5.279169649	85.9970374
		230-3A(138S)	0.45	0.043	4.1201245	0.52	0.044	4.652827487	12.9292329
		230-4A(138S)	0.34	0.044	3.0422334	0.42	0.044	3.758052971	23.5294118
138S	10/29/93	LAMINATE 300F/3 TONS							
		231-1A(TTS)	0.41	0.044	3.6685755	1.3	0.044	11.63206872	217.073171
		231-1A(TTS)	0.32	0.044	2.8832785	2.2	0.045	19.24759405	572.222222
		231-3A(138S)	0.49	0.044	4.3843951	0.68	0.044	6.084466714	38.7755102
		231-4A(138S)	0.52	0.044	4.6528275	0.65	0.044	5.816034359	25
232A	10/29/93	LAMINATE 300F/3 TONS							
		232-1A(TTS)	0.34	0.044	3.0422334				
		232-1A(TTS)	0.36	0.044	3.2211883				
		232-3A(138S)	0.57	0.044	5.1002147	0.71	0.044	6.352899069	24.5614035
		232-4A(138S)	0.58	0.044	5.1895922	0.62	0.044	5.547602004	6.89655172
233A	10/29/93	LAMINATE 300F/3 TONS							
		233-1A(TTS)	0.22	0.045	1.9247594				
		233-1A(TTS)	0.23	0.044	2.0575814				
		233-3A(138S)	0.28	0.044	2.5053686	2.25	0.044	20.13242663	703.571429
		233-4A(138S)	0.35	0.045	3.0621172	1.2	0.044	10.7372942	250.649351
234A	11/7/93	LAMINATE 300F/3 TONS							
		234-1A(TTS)	0.45	0.044	4.0264853				
		234-1A(TTS)	0.46	0.044	4.1159828				
		234-3A(138S)	0.5	0.044	4.4738126	1.05	0.044	9.395132427	110
		234-4A(138S)	0.64	0.044	5.7265569	1.35	0.043	12.36037356	115.843023
235A	11/7/93	LAMINATE 300F/3 TONS							
		235-1A(TTS)	0.46	0.044	4.1159828				
		235-1A(TTS)	0.44	0.044	3.937079				
		235-3A(138S)	0.76	0.044	6.8002663	0.66	0.044	5.905511811	-13.1578947
		235-4A(138S)	0.68	0.044	6.0844667	0.7	0.044	6.263421618	2.94117647
236A	11/7/93	LAMINATE 300F/3 TONS							
		236-1A(TTS)	0.68	0.033	8.1126223				

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237A	11/7/93	LAMINATE 300F/3 TONS							
		237-1A(1TTS)	0.67	0.043	6.1344076				
		237-1A(1TTS)	0.8	0.044	7.1581961				
		237-3A(138S)	0.67	0.043	6.1344076	0.96	0.044	8.589835361	40.027137
		237-4A(138S)	0.49	0.044	4.38443951	1.2	0.044	10.7372942	144.897959
238A	11/7/93	LAMINATE 300F/3 TONS							
		238-1A	0.42	0.044	3.758053	0.55	0.044	4.921259843	30.952381
		238-2A	0.38	0.045	3.3245844	0.4	0.045	3.499562555	5.26315789
		238-3A	0.36	0.044	3.22211883	0.48	0.044	4.294917681	33.3333333
		238-4A	0.34	0.045	2.9746282	0.5	0.045	4.374453193	47.0588235
239A	11/10/93	LAMINATE 300F/3 TONS							
		239-1A	0.459	0.045	4.015748	0.64	0.045	5.599300087	39.4335512
		239-2A	0.39	0.045	3.4120735	0.45	0.045	3.937007874	15.3846154
		239-3A	0.45	0.045	3.9370079	0.79	0.045	6.911636045	75.5555556
		239-4A	0.44	0.044	3.9370079	0.58	0.044	5.189692198	31.8181818
240A	11/16/93	LAMINATE 300F/3 TONS							
		240-1A	0.54	0.045	4.7244094	0.64	0.045	5.599300087	18.5185185
		240-2A	0.66	0.044	5.9055118				
		240-3A	0.6	0.045	5.2493438	0.84	0.044	7.516105941	43.1818182
		240-4A	0.77	0.044	6.8897638				
241A	11/15/93	LAMINATE 300F/3 TONS							
		241-1A	0.72	0.077	3.661358				
		241-2A	0.78	0.077	3.9881378	FOR BATTERY BUILD			
		241-3A	0.79	0.076	4.0924161				
242A	11/18/93	LAMINATE 300F/3 TONS							
		242-1A	0.59	0.066	3.5194464				
		242-2A	0.64	0.066	3.8177046	FOR BATTERY BUILD			
		242-3A	0.67	0.067	3.9370079				
		242-4A(NO PB)	0.72	0.066	4.2949177				
243A	11/18/93	LAMINATE 300F/3 TONS							
		243-1A	0.38	0.066	2.2667621				

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
244A	11/18/93	LAMINATE 300F/3 TONS 244-1A 244-2A 244-3A 244-4A(NO PB)	0.42 0.41 0.5 0.49	0.067 0.066 0.066 0.067	2.4679751 2.445717 2.9855817 2.4092138				
245A	12/1/93	LAMINATE 300F/3 TONS 245-1A 245-2A 245-3A 245-4A	0.59 0.55 0.66 0.7	0.041 0.042 0.041 0.041	5.6664504 5.1556055 6.3376224 6.7217208	0.62 0.71 0.77 1	0.041 0.041 0.041 0.04	5.953524102 6.817745243 7.393892837 9.842519685	5.08474576 32.2394678 16.666667 46.4285714
246A	12/13/93	LAMINATE 300F/3 TONS 246-1A 246-2A 246-3A 246-4A	0.6 0.42 0.54 0.42	0.019 0.018 0.019 0.019	12.432656 SAMPLE BROKE 9.1863517 11.189391 8.7028595	0.5 0.52 0.52 0.52	0.018 0.019 0.019 0.019	10.93613298 10.77496892 10.77496892	19.047619 -3.7037037 23.8095238
247A	12/13/93	LAMINATE 300F/3 TONS 247-1A 247-2A 247-3A 247-4A	0.285 0.34 0.36 0.31	0.02 0.02 0.02 0.02	5.6102362 6.6929134 7.0866142 6.1023622	0.39 0.38 0.52 0.45	0.02 0.021 0.02 0.02	7.677165354 7.124109486 10.23622047 8.858267717	36.8421053 6.44257703 44.4444444 45.1612903
248A	12/27/93	LAMINATE 300F/3 TONS 248-1A 248-2A 248-3A 248-4A	0.52 0.4 0.48 0.46	0.019 0.018 0.02 0.018	10.7744969 8.7489064 9.4488189 10.061242	1 0.66 1 0.66	0.019 0.018 0.02 0.019	20.72109407 14.43569554 19.68503937 13.67592209	92.3076923 65 108.333333 35.9267735
249A	1/5/94	LAMINATE 300F/3 TONS *249-1A 249-2A 249-3A 249-4A	0.88 0.38 0.38 0.4	0.02 0.019 0.019 0.02	17.322835 7.8740157 7.8740157 7.8740157	0.8 0.34 0.42 0.44	0.02 0.019 0.019 0.02	15.7480315 7.045171985 8.702859511 8.661417323	-9.09090909 -10.5263158 10.5263158 10
250A	1/5/94	LAMINATE 300F/3 TONS 250-1A 250-2A	0.88 0.5	0.041 0.041	8.4501632 4.8012291	0.84 0.46	0.041 0.041	8.066064913 4.417130785	-4.54545455 -8

*SAMPLE NOT SANDED PRIOR TO LAMINATION

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH)	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
251A	1/5/94	LAMINATE 300F/3 TONS 251-1A 251-2A	0.19 0.225	0.041 0.041	1.8244671 2.1605531	0.24 0.23	0.041 0.041	2.304589975 2.208565393	26.3157895 2.22222222
252A	1/7/94	LAMINATE 300F/3 TONS 252-1A 252-2A	0.15 0.125	0.041 0.042	1.4403687 1.1717285	0.195 0.18	0.041 0.042	1.872479355 1.687289089	30 4.4
253A	1/7/94	LAMINATE 300F/3 TONS 253-1A 253-2A(30 TONS)	0.15 0.155	0.019 0.011	3.1081641 5.547602	0.245 0.11	0.02 0.01	4.822834646 4.330708661	55.1666667 -21.9354839
254A	1/12/94	LAMINATE 300F/3 TONS 254-1A 254-2A 254-3A 254-4A	0.38 0.4 0.46 0.5	0.021 0.021 0.02 0.021	7.1241095 7.4990626 9.0551181 9.3738283	0.32 0.48 0.64 0.6	0.021 0.021 0.02 0.021	5.999250094 8.998875141 12.5984252 11.24859393	-15.7894737 20 39.1304348 20
255A	1/20/94	LAMINATE 300F 3 TONS/30 TONS 255-1A(3 TONS) 255-2A(3 TONS) 255-3A(30 TONS) 255-4A(30 TONS)	0.3 0.29 0.28 0.235	0.016 0.019 0.011 0.011	7.3818898 6.0091173 10.021475 8.4108805	0.265 0.28 0.32 0.295	0.016 0.019 0.011 0.011	6.520669291 5.801906341 11.45311382 10.5583393	-11.6666667 -3.448827586 14.2857143 25.5319149
256A	1/20/94	LAMINATE 300F/3 TONS NO SHIM 256-1A 256-2A 256-3A 256-4A	0.44 0.62 0.41 0.45	0.018 0.019 0.019 0.019	9.623797 12.847078 8.4956486 9.3244923	0.79 1.3 0.78 0.9	0.018 0.019 0.019 0.019	17.27909011 26.9374223 16.16245338 18.64898467	79.5454545 109.677419 90.2439024 100
257A	1/24/94	LAMINATE 300F/3 TONS .045", .031" SHIM 257-1A 257-2A 257-3A 257-4A 257-5A	0.76 0.74 0.8 0.9 0.45	0.04 0.04 0.04 0.041 0.04	7.480315 7.2834646 7.8740157 TAKE BATTERY #257 (12V) 8.6422124 FULL PB SHEET 7.5787402				

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE		RESISTANCE (OHM) AFTER		THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
			STABILITY TESTING	BEFORE	BEFORE	AFTER			
258A	1/25/94	LAMINATE 300F/3 TONS .045", .031" SHIM	258-1A	0.73	0.028	10.264342	0.91	0.028	12.795227559 24.6575342
			258-2A	0.79	0.028	11.107987	1.2	0.029	16.29106706 46.6608468
			258-3A	0.34	0.034	3.9370079	3.9370079	3.9370079	3.9370079
			258-4A	0.295	0.029	4.0048873	0.36	0.029	4.887320119 22.0338983
			258-5A	0.33	0.029	4.4800434	0.4	0.029	5.430355688 21.2121212
	1/26/94	LAMINATE 300F/3 TONS .045", .031" SHIM	259-1A	0.71	0.041	6.81177453			
			259-2A	0.78	0.043	7.1415492			
			259-3A	0.6	0.041	5.7614779	N UPON PRESSING IN THE PB SHEET		
			259-4A	0.69	0.04	6.7913386			
			259-5A	0.7	0.042	6.5616798			
259A	2/4/94	LAMINATE 300F/3 TONS .045", .031" SHIM	259-6A	0.7	0.029	9.5031225			
			259-7A	0.51	0.026	7.7225924	0.48	0.026	7.268322229 -5.88235294
			259-8A	0.51	0.027	7.4365704	0.54	0.027	7.874015748 5.88235294
			260-1A	0.56	0.041	5.3773766	DOUG- 4.5931759 TO MAKE 4V BATTERY 4.5931759 AMINATE BROKE	-FULL PB SHEET	
			260-2A	0.49	0.042	3.9370079			
			260-3A	0.35	0.03	5.511811			
	2/4/94	LAMINATE 300F/3 TONS .045", .031" SHIM	260-4A	0.46	0.028	6.4679415	0.54	0.027	7.874015748 21.7391304
			260-5A	0.34	0.026	5.1483949	0.49	0.026	7.419745609 44.1176471
			261A						
			261-1A	0.41	0.042	3.8432696	ULD NOT STICK TO LAMINATE		
			261-2A	0.42	0.042	3.9370079	"	"	
262A	2/4/94	LAMINATE 375F/3 TONS NO SHIM	261-3A	0.42	0.03	5.511811	"	"	
			261-4A	0.43	0.026	6.5112053	0.38	0.026	5.754088431 -11.627907
			261-5A	0.44	0.026	6.6626287	0.46	0.025	7.244094488 8.72727273
			262-1A	0.52	0.017	12.042612	0.74	0.016	18.20866142 51.2019231
			262-2A	0.6	0.017	13.895322	0.72	0.016	17.71653543 27.5
			262-3A	0.53	0.017	12.274201	0.69	0.016	16.97834646 38.3254717
			262-4A	0.51	0.017	11.811024	0.63	0.016	15.5019685 31.25

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH)	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
263A	2/1/94	LAMINATE 300F/3 TONS 0.045" SHIM	0.5	0.043	4.5779161 ATTERRY #263 6V-FULL PB SHEET				
		263-1A	0.49	0.043	4.4863578 "				
		263-2A							
264A	2/4/94	LAMINATE 300F/3 TONS 0.031" SHIM	0.46	0.034	5.3265401 MAKE BATTERY #264 4V				
		264-1A	0.41	0.034	4.7475683 STRATE CRACKED				
		264-2A							
265A	2/4/94	LAMINATE 300F/3 TONS 0.031" SHIM	0.3	0.033	3.5790981 MAKE BATTERY #265-6V				
		265-1A	0.28	0.033	3.3404915 "				
		265-2A							
		265-3A	0.33	0.032	4.0600394 DELAMINATED AT CORNER				
		265-4A	0.33	0.032	4.0600394 MAKE BATTERY #265-4V				
		265-5A	0.33	0.032	4.0600394 MINATE CRACKED				
266A	2/18/94	LAMINATE 300F/3 TONS 0.051" SHIM	0.36	0.046	3.0811366	0.37	0.045	3.237095363	5.0617284
		266-1A	0.38	0.045	3.3245844	0.48	0.044	4.294917681	29.1866029
		266-2A	0.41	0.046	3.5090722	0.84	0.045	7.349081365	109.430894
		266-3A							
		266-4A	0.4	0.045	3.4995626	0.46	0.045	4.024496938	15
267A	3-3-94	LAMINATE 300F/3 TONS							
		267-1A(C)	0.52	0.039	5.2493438	0.73	0.044	6.531853973	W/ PB SHEET
		267-2A(P)	0.61	0.037	6.4907427	0.6	0.041	5.761474938	"
		267-3A(P)	0.63	0.038	6.5271446	0.55	0.041	5.281352026	"
		267-4A(P)	0.45	0.036	4.9212598	0.56	0.041	5.377376608	"
		267-5A(P)	0.43	0.036	4.7025372	0.56	0.039	5.653139511	"
		267-6A(P)						5.31496063	INATED, PB SHEET
		267-7A(P)						4.921259843	"
		267-8A(C)						3.937007874	"
		267-9A(C)						3.641132283	"
		267-10A(C)						5.089302862	"
		267-11A(C)						5.377376608	"
		267-12A	0.45	0.036	4.9212598	0.44	0.037	4.681847202	-4.86486486
		267-13A	0.43	0.036	4.7025372	0.46	0.036	5.030621172	6.97674419
268A	3-3-94	LAMINATE 300F/3 TONS							
		268-1A	0.57	0.042	5.3430821	0.66	0.045	5.774278215	W/ PB SHEET

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
268A		268-2A	0.6	0.045	5.2493438	0.65	0.046	5.5631633	"
		268-3A	0.53	0.044	4.7423049	0.58	0.046	4.964053406	"
		268-4A	0.49	0.044	4.3843951	0.57	0.046	4.878466279	"
		268-5A	0.54	0.04	5.3149606	0.62	0.047	5.193499749	"
		268-6A				0.49	0.048	4.019028871	INATED, PB SH
		268-7A				0.46	0.048	3.772965879	"
		268-8A				0.5	0.049	4.017354973	"
		268-9A							
		MOVED FOR Q.C. DUE TO BLISTERING							
		268-10A							
		268-11A							
		268-12A	0.52	0.043	4.7610328	0.54	0.048	3.772965879	"
		268-13A	0.48	0.043	4.3947995	0.56	0.045	8.223972003	"
		LAMINATE 300F/3 TONS							
269A	3/3/94	269-1A	0.6	0.025	9.4488189	0.01	0.027	11.81102362	25
		269-2A	0.44	0.028	6.1867267	0.6	0.028	8.436445444	36.3636364
		269-3A	0.55	0.028	7.7334083	0.68	0.028	9.561304837	23.6363636
		269-4A	0.36	0.021	6.7491564	0.6	0.021	11.24859393	66.6666667
		LAMINATE 300F/3 TONS							
270A	3/4/94	270-1A	0.94	0.018	20.55993	0.35	0.018	7.655293088	-62.7659574
		270-2A	0.824	0.022	14.745884	0.98	0.022	17.53758053	18.9320388
		270-3A	0.745	0.02	14.665354	LASTIC CRACKED			
		270-4A	0.4	0.018	8.7488064	0.44	0.018	9.623797025	10
		270-5A	0.59	0.022	10.558339	0.74	0.021	13.87326584	31.3962873
		LAMINATE 300F/3 TONS							
271A	3/10/94	271-1A	0.37	0.02	7.2834646	0.39	0.02	7.677165354	5.40540541
		271-2A	0.34	0.02	6.6929134	0.31	0.02	6.102362205	-8.82352941
		271-3A	0.245	0.019	5.0766668	0.43	0.02	8.464566929	66.7346939
		271-4A	0.345	0.02	6.7913386	0.35	0.022	6.263421618	-7.77338603
		LAMINATE 300F/3 TONS							
272A	3/17/94	272-1A	0.3	0.028	4.2182227	0.58	0.027	8.457276174	100.493827
		272-2A	0.33	0.028	4.640045	0.65	0.028	9.139482556	96.969697
		272-3A	0.36	0.028	5.0618673	0.43	0.028	6.046119235	19.444444
		272-4A	0.36	0.028	5.0618673	0.41	0.028	5.764904387	13.8888889
		LAMINATE 300F/3 TONS							
273A	3/17/94	273-1A	0.38	0.04	3.7401575	0.94	0.041	9.026310736	141.335045
		273-2A	0.34	0.04	3.3464567	0.72	0.04	7.086614173	111.764706
		273-3A	0.39	0.041	3.7449987	0.65	0.041	6.241597849	66.6666667
		273-4A	0.37	0.042	3.4683165	0.52	0.041	4.993278279	43.9683586
		LAMINATE 300F/3 TONS							
274A	4/28/94	274-1A	2.8	0.052	21.199273	9	0.054	65.6167979	209.52381
		274-2A	3	0.058	20.363334	10	0.059	66.72894702	227.683616

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
275A	4/28/94	LAMINATE 300F/3 TONS 275-1A 275-2A	2.65 0.044 1.95 0.042	23.711525 50	24.5 0.044	0.044 0.044	219.2197566 447.3872584	824.528302 2347.55245	
276A	4/28/94	LAMINATE 300F/3 TONS 276-1A 276-2A	3.2 0.06 3.8 0.06	20.997375 24.934383	6.8 5	0.064 0.062	41.83070866 31.7500635	99.21875 27.3344635	
277A	5/2/94	LAMINATE 300F/3 TONS 277-1A 277-2A 277-3A 277-4A	0.58 NA NA NA	0.047 0.041 0.042 0.042	4.8584352 NA NA NA	DR 4V BATTERY 277-1C DR 4V BATTERY 277-2C IR 6V BATTERY 277-6V C " "			
278A	5/2/94	LAMINATE 300F/3 TONS 278-1A 278-2A 278-3A	NA NA NA	0.04 0.039 0.04	NA NA NA	DR 4V BATTERY 278-1C IR 6V BATTERY 278-6V C " "			
279A	5/2/94	LAMINATE 300F/3 TONS 279-1A 279-2A 279-3A 279-4A	NA NA NA NA	0.042 0.039 0.04 0.039	NA NA NA NA	DR 4V BATTERY 279-1C DR 4V BATTERY 279-2C IR 6V BATTERY 279-6V C " "			
280A	5/9/94	LAMINATE 300F/3 TONS 280-1A 280-2A 280-3A 280-4A	0.38 0.31 0.28 0.3	0.035 0.039 0.035 0.035	4.2744657 3.1294165 3.1496063 3.3745782	2.75 8.1 3 2.3	0.037 0.04 0.037 0.037	29.26154501 79.72440945 31.92168546 24.47329219	584.566145 2447.58065 913.513514 625.225225
281A	5/12/94	LAMINATE 300F/3 TONS 281-1A 281-2A	0.43 0.41	0.033 0.035	5.1300406 4.6119235	DR 4V BATTERY 281-1C DR 4V BATTERY 281-2C			

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM) BEFORE	THICKNESS (INCH) BEFORE	RESISTIVITY (OHM-CM) BEFORE	RESISTANCE (OHM) AFTER	THICKNESS (INCH) AFTER	RESISTIVITY (OHM-CM) AFTER	PERCENT CHANGE (%)
282A	5/13/94	LAMINATE 300F/3 TONS							
		282-1A	0.4	0.035	4.4994376 OR 4V BATTERY 282-1C				
		282-2A	0.42	0.037	4.469036 OR 4V BATTERY 282-2C				
		282-3A	0.38	0.036	4.1557305 OR 6V BATTERY 282-6V C				
		282-4A	0.4	0.036	4.3744532 " " "				
283A	5/13/94	LAMINATE 300F/3 TONS							
		283-1A	0.52	0.025	8.1889764	2.85	0.025	44.88186976	448.076923
		283-2A	0.5	0.025	7.8740157	2.35	0.025	37.00787402	370
		283-3A	0.45	0.025	7.0866142	3.3	0.025	51.96850394	633.333333
		283-4A	0.36	0.024	5.9055118	2.5	0.024	41.01049869	594.444444
284A	5/25/94	LAMINATE 300F/3 TONS							
		284-1A	0.5	0.041	4.8012291	1.1	0.041	10.56270405	120
		284-2A	0.54	0.041	5.1653274	1.4	0.041	13.44344152	159.259259
		284-3A	0.55	0.041	5.281352	1.6	0.041	15.36393317	190.909091
		284-4A	0.7	0.04	6.8897638	1.6	0.041	15.36393317	122.996516
285A	6/2/94	LAMINATE 300F/3 TONS							
		285-1A	0.89	0.045	7.7865267 OR 4V BATTERY 285-1				
		285-2A	1.15	0.046	9.8425197				
		285-3A	1.25	0.048	10.255625				
		285-4A	1.35	0.047	11.308427				
286A	6/2/94	LAMINATE 300F/3 TONS							
		286-1A	1.1	0.047	9.2142737 DONT USE				
		286-2A	1.25	0.049	10.043387 OR 4V BATTERY 286-2				
		286-3A	1.05	0.05	8.2677165				
		286-4A	1.25	0.051	9.6495291 DONT USE				
287A	6/3/94	LAMINATE 300F/3 TONS							
		287-1A	0.9	0.047	7.5389512 DONT USE				
		287-2A	0.6	0.043	5.4934394 OR 4V BATTERY 287-2				
		287-3A	0.595	0.044	5.3239084 OR 4V BATTERY 287-3				
		287-4A	0.53	0.043	4.8525911				
288A	6/15/94	LAMINATE 300F/3 TONS							
		288-1A	0.66	0.041	6.3376224 E, SUBSTRATE CRACKED				

SAMPLE NUMBER	TEST DATE	MATERIAL COMPOSITION	RESISTANCE (OHM)		THICKNESS (INCH)		RESISTANCE (OHM-CM)		THICKNESS (INCH)		RESISTIVITY (OHM-CM)		PERCENT CHANGE (%)
			BEFORE	BEFORE	BEFORE	BEFORE	AFTER	AFTER	AFTER	AFTER	AFTER	AFTER	
288A	6/16/94	LAMINATE 300F/3 TONS	288-2A	0.8	0.042	7.4990626	OR 4V BATTERY 288-2						
			288-3A	0.54	0.041	5.1853274							
			288-4A	0.92	0.042	8.623922							
289A	6/23/94	LAMINATE 300F/3 TONS	289-1A	0.56	0.04	5.5118111	OR 4V BATTERY 289-1						
			289-2A	0.52	0.04	5.1181102							
			289-3A	0.53	0.041	5.0893029							
			289-4A	0.55	0.04	5.4133858							
290A	6/23/94	LAMINATE 300F/3 TONS	290-1A	0.68	0.018	14.873141	OR 4V BATTERY 290-1						
			290-2A	0.7	0.019	14.504766	OR 4V BATTERY 290-6V						
			290-3A	0.62	0.019	12.847078	"						
			290-4A	0.66	0.02	12.992126	"						
			290-5A	0.52	0.02	10.23622							
			290-6A	0.49	0.02	9.6456693							
			290-7A	0.44	0.019	9.1172814							
			290-8A	0.5	0.019	10.3860547							
			290-9A	0.5	0.021	9.3738283							
			290-10A	0.5	0.021	9.3738283							
			290-11A	0.52	0.021	9.7487814							
			290-12A	0.54	0.021	10.123735							

APPENDIX B

DELIVERABLE DATA

BUILD ID

WPG-6

Description

12 V Bipolar Battery

ASSEMBLY

Substrate Type	5.9375" X 9.1875" X 0.012" tin-lead alloy sheet							
Grid Type	0.016" thick metallic screen soldered to the substrate							
Separator Type, Dimensions	5.125" X 8.562" X 0.029"							
Positive Paste Density	3.35 g/cc							
Negative Paste Density	3.75 g/cc							

Plate ID	PTE D2	D5		D7		D8		D9		D10		NTE D4
Pb Mass (g.)	260.90	158.80		160.20		162.60		158.10		161.60		261.90
AM Mass (g.)	51.70	104.30		104.20		106.00		103.50		104.80		53.40
Dry AM (g.)	51.70	52.19	52.11	52.52	51.68	52.92	53.08	52.26	51.24	51.94	52.86	53.40
Sep. Mass (g.)	Cell 1	3.54	Cell 2	3.52	Cell 3	3.53	Cell 4	3.53	Cell 5	3.52	Cell 6	3.51

Termination	Copper stud soldered to terminal electrode							
Containment Type	Solvent bonded ABS. Container core thickness = 0.668"							
Completed Mass	3.5121 kg							

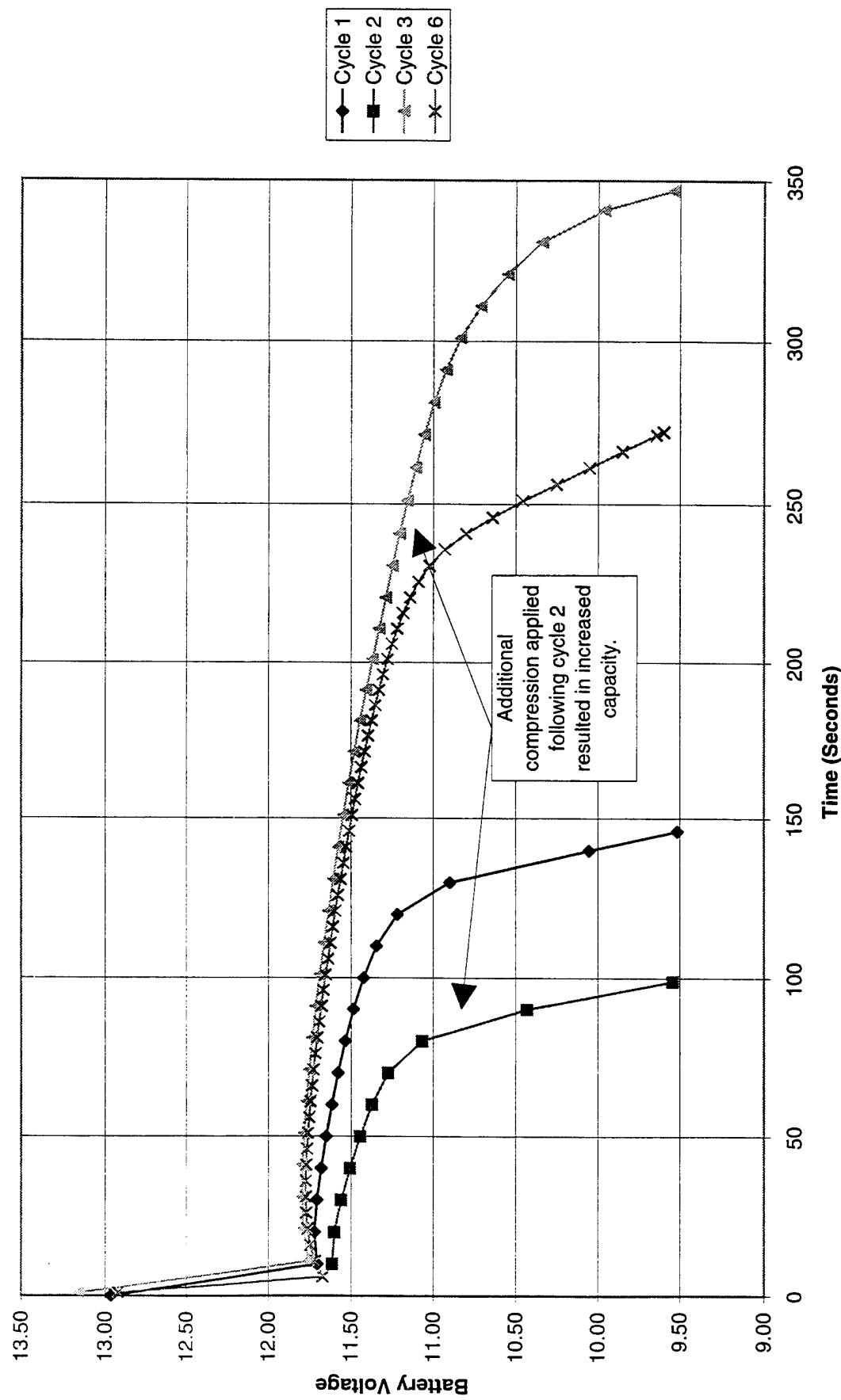
FORMATION

Acid Gravity	Chilled 1.265							
% Sodium Sulfate	1.5							
Method of Fill	Vacuum							
Time	27H:55M:04S							
Amps	1.0							
Voltage Limit	16.32							
Amp Hours	20.62							
Watt Hours	311.8							
Internal Resistance	13.5 mΩ							

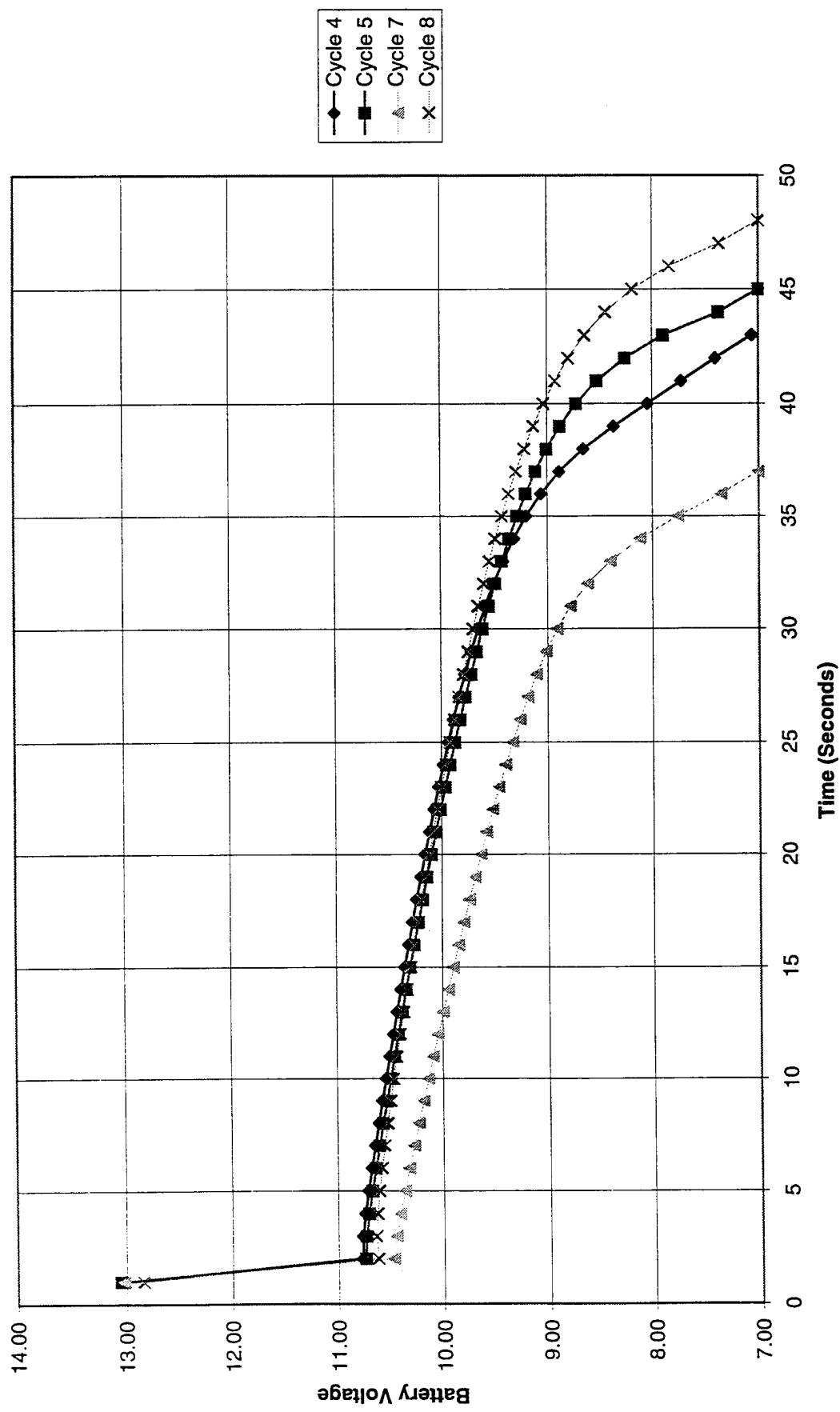
CYCLING HISTORY

Cycle	Date	IR (mW)	OCV	Discharge				Recharge				
				Amps	EODV	Ah	Wh	Amps	Vlimit	Ah	Wh	% Rchg
1	11/15/95	13.5	12.966	21	9.6	0.85	9.6	0.5	15.30	0.935	12.92	110
2	11/16/95	16.5	NA	21	9.6	0.57	6.4	0.1	14.40	NA	NA	NA
3	11/20/95	10.5	13.158	21	9.6	2.01	22.8	0.5	14.40	2.211	29.48	110
4	11/21/95	8.2	13.019	124	7.2	1.44	14.1	0.5	14.40	1.584	21.22	110
5	11/22/95	8.6	13.05	124	7.2	1.51	14.7	0.5	14.40	1.661	22.24	110
6	11/30/95	10.0	12.922	21	9.6	1.58	17.9	0.5	14.40	1.738	23.20	110
7	12/1/95	9.8	13.017	124	7.2	1.23	11.7	0.5	14.40	1.353	18.12	110
8	12/11/95	8.8	12.84	124	7.2	1.61	15.6	0.5	14.40	1.771	23.42	110

WPG-6
21 Amp Discharge Curves



WPG-6
124 Amp Discharge Curves



BUILD ID WPG-8

Description 24 V Bipolar Battery

ASSEMBLY

Substrate Type	5.9375" X 9.1875" X 0.012" tin-lead alloy sheet									
Grid Type	0.016" thick metallic screen soldered to the substrate									
Separator Type, Dimensions	5.125" X 8.562" X 0.029"									
Positive Paste Density	3.51 g/cc									
Negative Paste Density	3.83 g/cc									

Plate ID	PTE D54	D14		D15		D17		D18		D20		D21	
Pb Mass	258.70	162.90		162.20		161.90		162.80		163.10		162.00	
AM Mass	52.10	106.00		105.30		104.70		104.80		105.40		103.60	
Dry AM	52.10	52.71	53.29	52.41	52.89	52.65	52.05	52.33	52.47	52.37	53.03	51.85	51.75
Sep. Mass	Cell 1	3.52	Cell 2	3.53	Cell 3	3.48	Cell 4	3.52	Cell 5	3.48	Cell 6	3.47	Cell 7

Plate ID	D22		D23		D25		D26		D27		NTE D57	
Pb Mass	160.40		163.10		160.90		161.90		162.80		258.50	
AM Mass	103.20		102.00		106.00		101.70		103.30		54.00	
Dry AM	52.05	51.15	51.24	50.76	51.22	54.78	50.75	50.95	51.51	51.79	54.00	
Sep. Mass	3.49	Cell 8	3.46	Cell 9	3.49	Cell 10	3.50	Cell 11	3.51	Cell 12	3.50	

Termination	Copper stud soldered to terminal electrodes											
Containment Type	Solvent bonded ABS. Container core thickness = 1.153".											
Containment Mass	5.5360 kg											

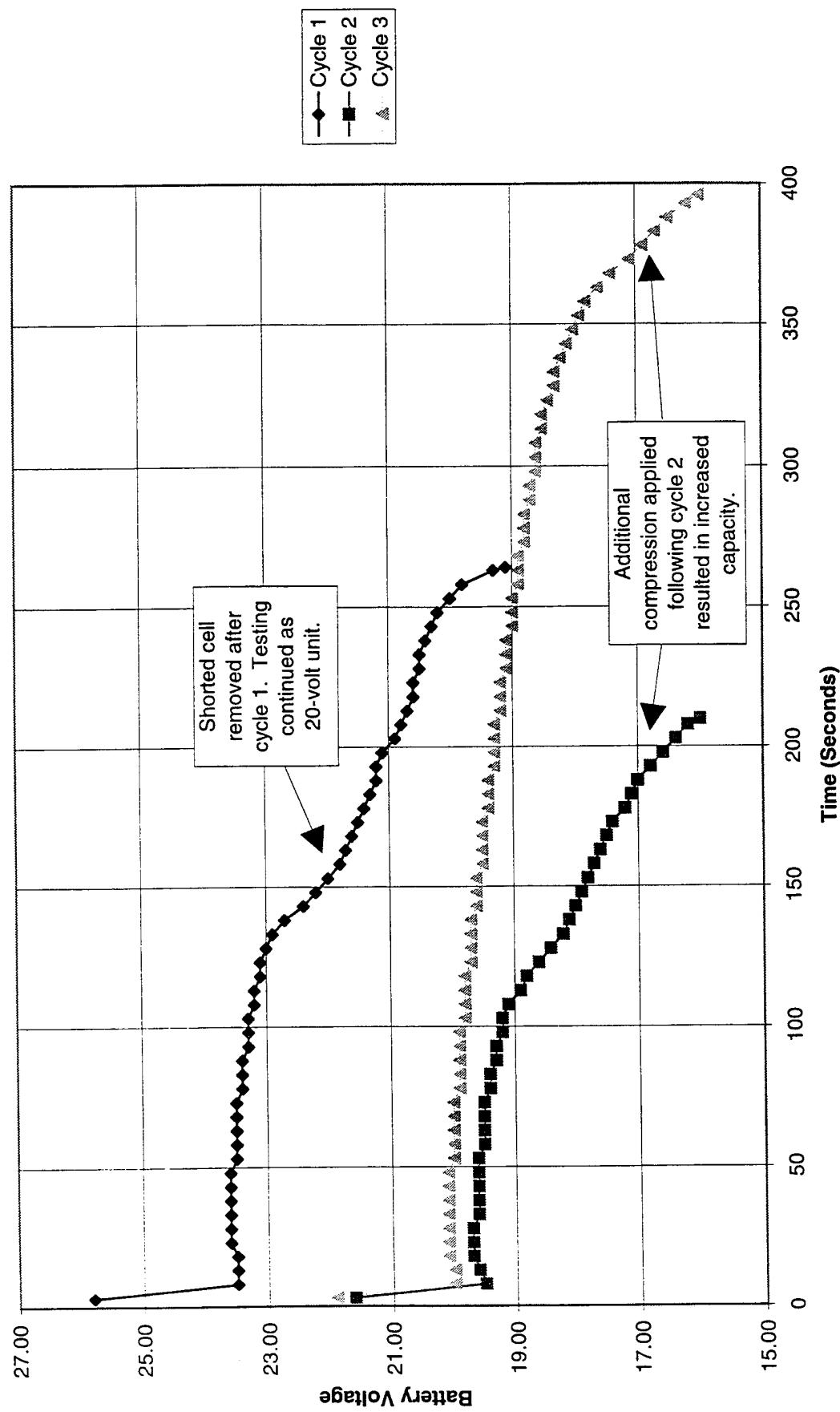
FORMATION

Acid Gravity	Chilled 1.265											
% Sodium Sulfate	1.5											
Method of Fill	Vacuum											
Time	20H:37M:03S											
Amps	1.0											
Voltage Limit	32.64											
Amp Hours	20.62											
Watt Hours	594.0											
Internal Resistance	14.0 mΩ											

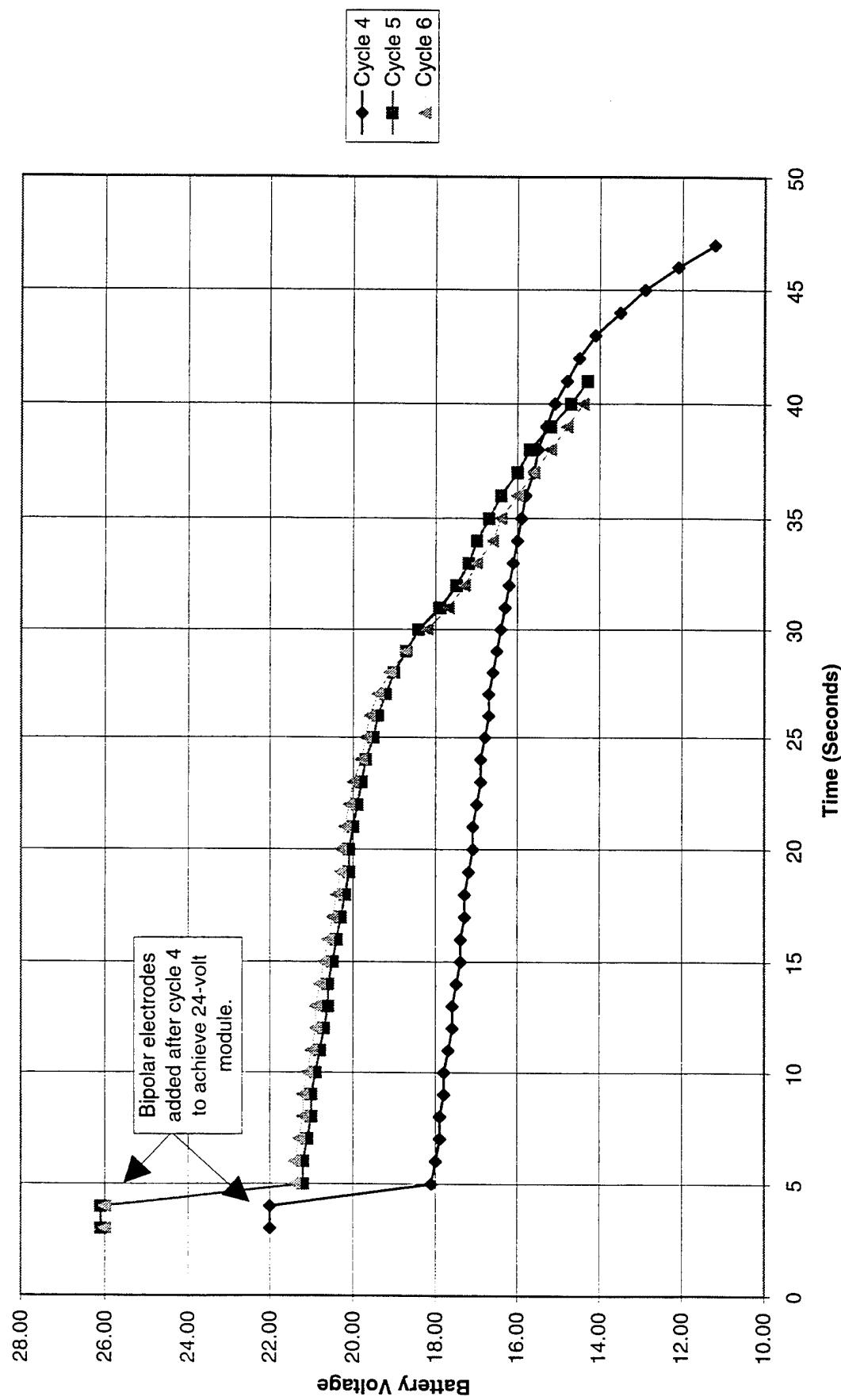
CYCLING HISTORY

Cycle	Date	IR (mW)	OCV	Discharge				Recharge				
				Amps	EODV	Ah	Wh	Amps	Vlimit	Ah	Wh	% Rchg
1	1/16/96	14.5	25.80	21	19.2	1.50	31	0.5	30.60	1.65	44	110
1/16/96 Two shorted bipolar electrodes removed. Continue cycling as 20-volt nominal battery.												
2	1/18/96	17.5	21.60	21	16.0	1.20	20	1.0	25.50	1.32	29	110
3	1/18/96	12.5	21.90	21	16.0	2.29	41	0.1	25.50	2.51	50	110
4	1/19/96	12.5	22.00	124	12.0	1.48	23	0.1	25.50	1.62	32	110
1/23/96 Two good bipolar electrodes added to stack to achieve 24-volt module.												
5	1/24/96	17.0	26.10	124	14.4	1.27	23	0.1	30.60	1.39	28	110
6	1/26/96	16.0	26.00	124	14.4	1.24	23	0.1	30.60	1.36	27	110

WPG-8
21 Amp Discharge Curves



WPG-8
124 Amp Discharge Curves



BUILD ID

WPG-11

Description

12 V Bipolar Battery

ASSEMBLY

Substrate Type	5.9375" X 9.1875" X 0.012" tin-lead alloy sheet						
Grid Type	0.016" thick metallic screen soldered to the substrate						
Separator Type, Dimensions	5.125" X 8.562" X 0.029"						
Positive Paste Density	3.40 g/cc						
Negative Paste Density	3.75 g/cc						

Plate ID	PTE D72	D66		D57		D69		D64		D65		NTE D74
Pb Mass (g.)	261.03	160.07		160.71		163.42		163.13		164.39		258.98
AM Mass (g.)	50.97		102.23		102.49		102.98		101.27		101.91	54.32
Dry AM (g.)	50.97	51.03	51.20	50.97	51.52	51.23	51.75	50.40	50.87	50.57	51.34	54.32
Sep. Mass (g.)	Cell 1	3.53	Cell 2	3.45	Cell 3	3.48	Cell 4	3.52	Cell 5	3.50	Cell 6	3.48

Termination	Copper stud soldered to terminal electrode						
Containment Type	Solvent bonded ABS. Container core thickness = 0.671".						
Containment Mass	3.4908 kg						

FORMATION

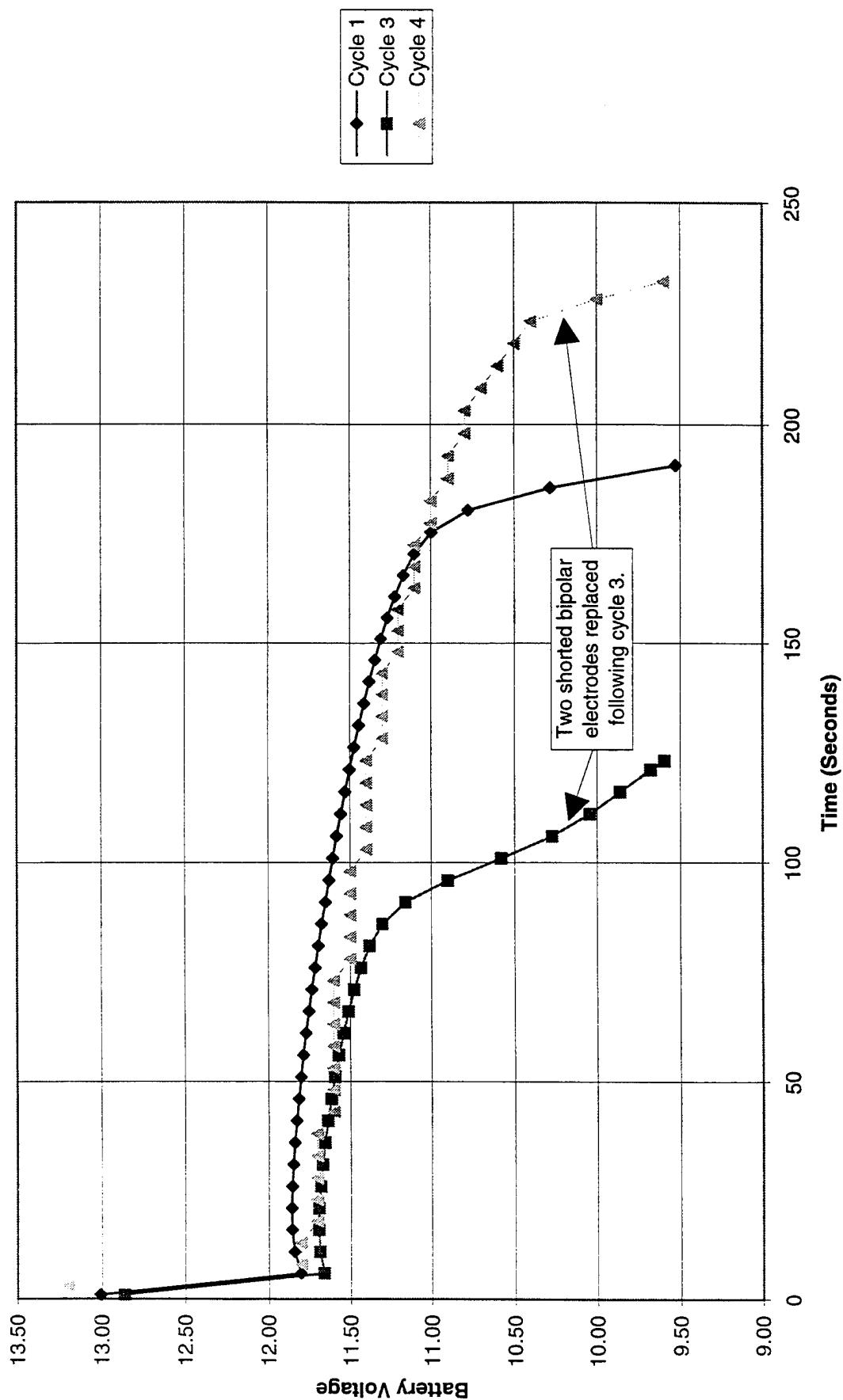
Acid Gravity	Chilled 1.265						
% Sodium Sulfate	1.5						
Method of Fill	Vacuum						

Time							
Amps	1						
Voltage Limit	16.32						
Amp Hours	20.62						
Watt Hours	NA						
Internal Resistance	12 mΩ						

CYCLING HISTORY

Cycle	Date	IR (mW)	OCV	Discharge				Recharge				
				Amps	EODV	Ah	Wh	Amps	Vlimit	Ah	Wh	% Rchg
1	2/16/96	10.5	13.009	21	9.6	1.1	12.7	0.5	15.30	1.21	16.4	110
2	2/16/96	11.0	13.137	124	7.2	0.72	6.6	0.5	15.30	0.79	10.8	110
3	2/19/96	12.0	12.866	21	9.6	0.71	7.9	0.5	15.30	0.78	10.5	110
2/26/96 Replaced two shorted bipolar electrodes.												
4	2/27/96	11.5	13.200	21	9.6	1.33	12.0	0.5	14.40	1.46	17.0	110
5	2/28/96	11.0	13.005	124	7.2	0.82	7.0	0.5	14.40	0.90	10.0	110

WPG-11
21 Amp Discharge Curves



WPG-11
124 Amp Discharge Curves

